

BMJ Open is committed to open peer review. As part of this commitment we make the peer review history of every article we publish publicly available.

When an article is published we post the peer reviewers' comments and the authors' responses online. We also post the versions of the paper that were used during peer review. These are the versions that the peer review comments apply to.

The versions of the paper that follow are the versions that were submitted during the peer review process. They are not the versions of record or the final published versions. They should not be cited or distributed as the published version of this manuscript.

BMJ Open is an open access journal and the full, final, typeset and author-corrected version of record of the manuscript is available on our site with no access controls, subscription charges or pay-per-view fees (http://bmjopen.bmj.com).

If you have any questions on BMJ Open's open peer review process please email info.bmjopen@bmj.com

# **BMJ Open**

# Adverse Childhood Experiences and Diabetes in Adulthood: Results of the Singapore Mental Health Study

Journal:	BMJ Open
Manuscript ID	bmjopen-2020-045167
Article Type:	Original research
Date Submitted by the Author:	24-Sep-2020
Complete List of Authors:	Subramaniam, Mythily; Institute of Mental Health, Research Division Abdin, Edimansyah; Institute of Mental Health, Research Division Vaingankar, Janhavi; Institute of Mental Health Singapore, Research Chang, Sherilyn; Institute of Mental Health, Research Division Sambasivam, Rajeswari Jeyagurunathan, Anitha; Institute of Mental Health, Singapore, Research Seow, Lee Seng Esmond; Institute of Mental Health, Research Division Van Dam, Rob; National University Singapore Saw Swee Hock School of Public Health Chow, Wai Leng; Government of Singapore Ministry of Health Chong, Siow Ann; Institute of Mental Health,
Keywords:	General diabetes < DIABETES & ENDOCRINOLOGY, EPIDEMIOLOGY, Child protection < PAEDIATRICS

SCHOLARONE™ Manuscripts



I, the Submitting Author has the right to grant and does grant on behalf of all authors of the Work (as defined in the below author licence), an exclusive licence and/or a non-exclusive licence for contributions from authors who are: i) UK Crown employees; ii) where BMJ has agreed a CC-BY licence shall apply, and/or iii) in accordance with the terms applicable for US Federal Government officers or employees acting as part of their official duties; on a worldwide, perpetual, irrevocable, royalty-free basis to BMJ Publishing Group Ltd ("BMJ") its licensees and where the relevant Journal is co-owned by BMJ to the co-owners of the Journal, to publish the Work in this journal and any other BMJ products and to exploit all rights, as set out in our licence.

The Submitting Author accepts and understands that any supply made under these terms is made by BMJ to the Submitting Author unless you are acting as an employee on behalf of your employer or a postgraduate student of an affiliated institution which is paying any applicable article publishing charge ("APC") for Open Access articles. Where the Submitting Author wishes to make the Work available on an Open Access basis (and intends to pay the relevant APC), the terms of reuse of such Open Access shall be governed by a Creative Commons licence – details of these licences and which Creative Commons licence will apply to this Work are set out in our licence referred to above.

Other than as permitted in any relevant BMJ Author's Self Archiving Policies, I confirm this Work has not been accepted for publication elsewhere, is not being considered for publication elsewhere and does not duplicate material already published. I confirm all authors consent to publication of this Work and authorise the granting of this licence.

Title: Adverse Childhood Experiences and Diabetes in Adulthood: Results of the Singapore Mental Health Study

Authors: Mythily Subramaniam,<sup>1,2</sup> Edimansyah Abdin,<sup>1</sup> Janhavi Vaingankar,<sup>1</sup> Sherilyn

Chang, <sup>1</sup> Rajeswari Sambasivam, <sup>1</sup> Anitha Jeyagurunathan, <sup>1</sup> Esmond Seow, <sup>1</sup> Rob M. van

Dam,<sup>2</sup> Wai Leng Chow,<sup>3</sup> Siow Ann Chong<sup>1</sup>

<sup>1</sup>Research Division

**Institute of Mental Health** 

**Singapore** 

<sup>2</sup> Saw Swee Hock Public Health

**Singapore** 

<sup>3</sup> Epidemiology and Communicable Diseases Division

**Ministry Of Health** 

**Singapore** 

**Correspondence:** 

**Mythily Subramaniam** 

**Director, Research Division** 

**Institute of Mental Health** 

**Buangkok Green Medical Park** 

10 Buangkok View

Singapore 539747

Tel: 65-6389 3633 Fax: 65-6343 7962

Email: Mythily@imh.com.sg

Word count: 4000



#### Abstract

Objectives Several studies have linked diabetes mellitus to adverse childhood experiences (ACE) While a number of studies have examined the association between ACEs and diabetes in Western populations, few have done it in Asian populations. The current study aimed to examine (i) the association between ACEs and diabetes, including the association after age stratification and (ii) the association of comorbid depression, resource utilization and health related quality of life (HRQoL) among those with diabetes and ACEs in Singapore.

**Settings** Participants were surveyed in their homes or any other preferred venue of their choice. **Participants** 6126 individuals aged 18 years and above randomly selected among Singapore residents.

**Design** Cross-sectional nationwide epidemiological study.

**Results** Exposure to any ACE was not associated with increased odds of diabetes, however those who had experienced parental separation, death or divorce of a parent, had higher odds of diabetes. In addition, we observed significant interaction between age and ACEs in relation to odds of diabetes. ACEs were significantly associated with higher odds of diabetes mainly in the younger age group. The prevalence of major depressive disorder was significantly higher among those with diabetes and ACEs than those with diabetes alone (3.7% and 0.3% respectively). **Conclusion** Efforts to promote regular exercise and healthy lifestyles both in the population and among those with diabetes, must continue for the prevention and management of diabetes. The findings emphasise the need to create more awareness of both the prevalence and impact of ACEs among those treating chronic diseases.

Key words: Adverse childhood experiences; Diabetes; Depression; Healthcare utilization; Asian

# **Article Summary**

# Strengths and limitations of this study

- 1. The study reports on the association of Adverse Childhood Events (ACEs) with diabetes based on data from a large representative sample of the Singapore population.
- 2. Data on ACEs was captured retrospectively and is there is the possibility of recall bias and given the sensitivity of the questions there could have been a tendency to underreport the experience.
- 3. The study did not assess the severity, age of onset and duration of ACEs which may have an impact on the development of diabetes.
- 4. The cross-sectional design limits assertions regarding the sequence of causal events, but this was overcome to a significant extent by excluding those with onset of diabetes before 18 years of age.

#### Introduction

Adverse Childhood Experiences (ACEs) is a broad construct that includes neglect, abuse as well as household dysfunction resulting from parental mental illness, substance abuse, imprisonment, domestic violence and divorce occurring before a child reaches 18 years of age. The prevalence of ACEs varies considerably across studies depending on the population and the methodology. A study from the USA found an overall prevalence of 46% while another study estimated that about 47% of the population of England had experienced at least one such ACE3 ACEs impact health outcomes across the life course. These include emergence of mental and physical illnesses, 5-7 premature death and higher health care utilization. Studies have attributed the higher utilization among those with ACEs to health comorbidities, functional disability, need for complex care coordination and frequent missing of appointments.

Diabetes mellitus (subsequently referred to as diabetes) is a chronic metabolic condition with an estimated global prevalence of 9.3% (463 million people) in 2019, which is projected to increase to 10.2% (578 million) by 2030.<sup>14</sup> The annual direct cost of diabetes—i.e., cost of treating and managing the disease and its complications, was calculated to be International\$ 825 billion in 2014.<sup>15</sup> Risk factors commonly identified for type 1 diabetes include, genetic susceptibility, viral infections, and toxins while type 2 diabetes has been strongly linked with increasing age, increased body mass index (BMI), ethnicity, and family history.<sup>16</sup>

Over the years, studies have linked diabetes to ACEs. A systematic review and metaanalysis suggested that adverse childhood experiences (ACEs) were associated with a significantly elevated risk of type 2 diabetes in adulthood. Participants who had experienced an ACE had a 32% increased risk of developing type 2 diabetes later in life. Additionally, they found a differential effect of the type of ACE; neglect had the strongest effect while physical abuse had the least.<sup>17</sup> Studies have further shown that endorsement of a single ACE may increase risk for diabetes in adulthood by 11% and this risk may increase two-fold when there are more than three ACEs. 19

Suggested pathophysiological mechanisms linking ACEs with diabetes include, effects on the hypothalamic- pituitary axis, <sup>20-22</sup> metabolic dysregulation i.e., elevated level of triglycerides, free fatty acids, glucose, and insulin caused by the chronic stress from ACEs, <sup>23</sup> elevated levels of inflammatory markers, <sup>24</sup> and epigenetic changes. <sup>25</sup> <sup>26</sup> Psycho-social risk factors have also been proposed as a possible mechanism. Those with a history of ACEs are more likely to have health-risk behaviors such as cigarette smoking, alcohol misuse, infrequent physical activity and having higher BMIs<sup>27-29</sup> - all of which can increase the risk of diabetes. Lastly, studies have shown a significant association of ACEs with posttraumatic stress disorder (PTSD) and depression which in turn are strongly associated with the development of obesity, diabetes and cardiovascular diseases. <sup>19</sup> <sup>23</sup> <sup>30</sup> However, much less is known about the correlates of ACEs for adults who have diabetes. Although in general exposure to ACEs is associated with increased burden to physical illness such as depression, <sup>31</sup> and lower health-related quality of life (QOL), <sup>32</sup> it is not known how or if these variables are related to ACEs in patients who have diabetes.

Singapore is a multi-ethnic country located in Southeast Asia. A recent study found that the lifetime prevalence of ACEs in the adult population of Singapore was 63.9%. ACEs were associated with several mental disorders including major depressive disorder (MDD) and alcohol use disorder.<sup>33</sup> However, the study did not examine the association of ACEs with physical conditions. The prevalence of diabetes in Singapore is higher compared to the global prevalence, with about 1 in 9 Singaporeans (11.3%) suffering from the disease.<sup>34</sup> Diabetes was the seventh leading cause of disability adjusted life years in Singapore in 2017.<sup>35</sup> While a number of studies have examined the association between ACEs and diabetes in Western populations, few have done

it in Asian populations. It is also not known whether those with diabetes and ACEs are more likely to be associated with mental disorders or increased resource utilization.

Accordingly, the current study aimed to examine (i) the association between ACEs and diabetes, including the association after age stratification and (ii) the association of comorbid depression, resource utilization and health related quality of life (HRQoL) among those with diabetes and ACEs. The authors hypothesized that experiencing ACEs in childhood would be associated with diabetes, and that the association will be attenuated by health risk behaviors, including obesity, smoking and alcohol use. We also hypothesized that those with diabetes and ACEs would be significantly associated with MDD, increased resource utilization and lower HRQoL.

#### Methods

The current study comprises secondary analysis of data from the Singapore Mental Health Study conducted in 2016 (SMHS 2016). SMHS 2016 was a population-based, psychiatric epidemiological study conducted among Singapore residents. The study has been described in detail in an earlier article.<sup>36</sup> In brief, a probability sample was randomly selected using a disproportionate stratified sampling design from a national registry of Singaporeans and permanent residents. Residents aged 65 and above, Malays and Indians were over-sampled to ensure that an adequate sample size would be achieved to improve the reliability of estimates for the subgroup analysis. In all, 6,126 respondents were interviewed as part of the study with a response rate of 69.5%.<sup>36</sup>

#### 2.1. Study procedures

Face to face interviews were conducted in English, Chinese or Malay language depending on the preference of the respondents. Respondents comprised those who were 18 years and older,

Singapore citizens or Permanent residents, living in Singapore in a community dwelling during the field period. Trained interviewers from a survey research company administered the survey using computer assisted personal interviews (CAPI). The study was approved by the National Healthcare Group's Domain Specific Review Board. Written informed consent was obtained from all participants and parents or legally acceptable representatives of those aged below 21 years.

## 2.2. Questionnaires

2.2.1. World Health Organisation – Composite International Diagnostic Interview version 3.0 (WHO-CIDI 3.0)

Diagnoses of lifetime MDD, was generated using the CIDI 3.0 algorithm for the Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV) criteria.<sup>37</sup>

A modified version of the CIDI checklist of chronic medical disorders was used to establish the prevalence of eighteen chronic medical conditions including diabetes.<sup>38</sup> The question was read as 'I'm going to read to you a list of health problems some people have. Has a doctor ever told you that you have any of the following...' This was followed by a list of chronic medical conditions which were considered prevalent in Singapore's population. Those who gave a positive answer to the question were routed to the question "How old were you when you were first diagnosed with the condition?"

# 2.2.2. Adverse Childhood Experiences—International Questionnaire (ACE -IQ)

The World Health Organisation's international questionnaire was used to establish the prevalence of ACEs in the current study.<sup>39</sup> The questions from ACE-IQ included in the current study cover physical, sexual and emotional abuse and neglect by parents or caregivers; as well as family dysfunction. Only one question was asked to assess sexual abuse – "Did someone touch or fondle you in a sexual way when you did not want them to?" All questions about ACEs pertained to the respondents' first 18 years of life, and responses were binary (yes or no) or frequency-based.

From these, a dichotomous variable was created to reflect exposure to each ACE type and category (abuse, neglect, household dysfunction, and bullying). Given the sensitive nature of the questions, the questionnaire was self-administered by the respondent using a tablet. For those who were not literate, the items were read out in a neutral manner by the interviewer. Consistent with previous research,<sup>40-42</sup> the number of ACEs that the respondents reported having experienced was summed into an ACE count (range 0 to 11) and categorised into four groups for analysis: 0 ACE, 1 ACE, 2 ACEs and 3 + ACEs.

### 2.2.3 Health Risk Behaviors

- *Overweight and obesity:* BMI was calculated as weight in kilograms divided by the square of height in meters; weight and height were self-reported by participants. BMI was coded into underweight (< 18.5), normal (18.5–24.9), overweight (25–29.9), and obese (> = 30), consistent with current recommendations.<sup>43</sup>
- *Alcohol use*: Based on questions from the WHO-CIDI 3.0 as well as a series of questions created to assess binge drinking, participants were classified as non-drinkers, drinker but not binge drinker and those with binge drinking.
- *Smoking history:* Participants were asked if they are current smokers, ex-smokers or non-smokers i.e., have never smoked before. They were then classified as never, past, and current smokers.

# 2.2.3. SF-12 questionnaire

The Short Form (SF)-12 instrument was used to measure the generic HRQoL.<sup>44</sup> It measures eight domains of health: physical function (PF), role-physical (RP), bodily pain (BP), general health (GH), vitality (VT), social function (SF), role-emotion (RE), and mental health (MH). Scores from these domains are used to generate two summary scores - physical component summary (PCS)

and mental component summary (MCS). Both summary scores range between 0 and 100, with higher scores indicating better health. Internal consistency reliability for PCS (Cronbach's alpha = 0.81) and MCS (Cronbach's alpha = 0.82) domain scores were high in the current study.

#### 2.2.4 Resource utilization

Information on healthcare resource utilization was obtained from respondents using an adapted version of the Client Service Receipt Inventory (CSRI).<sup>45</sup> The instrument asked whether respondents had accessed specific healthcare resources during the past 3-month period before the time of the interview. This included care from the polyclinic doctor (government primary care doctors), restructured hospital doctors (a public hospital doctor which is wholly owned by the government), private hospital/clinic doctors, and hospitalisations.

## 2.2.5 Socio-Demographic Questionnaire

Data on sex, age, ethnicity (Chinese, Malay, Indian, and Others), marital status (not married, married, divorced/ separated and widowed), educational level (primary and below, secondary, vocational/ institute of technical education (ITE), pre-university/ junior college, diploma and university), employment status (employed, unemployed and economically inactive, i.e., students, homemakers and retirees) and monthly household income was collected. Monthly household income was calculated as the average pre-tax income in the past 12 months, of all family members living in the same household.

#### Statistical analysis

Statistical analyses were carried out using the STATA version 13.0. All estimates were analyzed using survey weights to adjust for oversampling, non-response and poststratification

according to age and ethnicity distribution between the survey sample and the Singapore adult population in 2014. Mean and standard errors were calculated for continuous variables and frequencies and percentages for categorical variables. Multiple logistic regression was used to determine sociodemographic factors associated with diabetes as well as to determine the association of ACEs (specific experiences and counts) with diabetes (model 1), after adjusting for health risk behavior factors (model 2) and adjusting for both health risk behaviors and sociodemographic factors (model 3). In addition, we tested for interaction between age (18-49 years, and 50+ yrs) and different ACEs (yes/no) in relation to diabetes by including multiplicative interaction terms in the multivariable logistic regression models. We chose the cut-off age as 49 years as the median age of onset of diabetes in the sample was 49 years. Since the interaction between age and several ACEs was significant, we described the associations between ACEs and diabetes stratified for age.

Multiple linear and logistic regression models were also used to examine the associations among those with diabetes and any ACE (versus among those with diabetes alone) and MDD, primary care visits, restructured hospital doctor visits, private hospital doctor visits, and number of hospitalisations after adjusting for health risk behavior factors and sociodemographic factors.

Those with onset of diabetes up to the age of 18 years were excluded from these analyses as ACEs measure adversities experienced until 18 years. All statistically significant differences were evaluated at the 0.05 level using 2-sided tests.

## Patient and public involvement

Patients or participants of this study were not involved in the design or development of the study.

#### Results

Socio-demographic characteristics of the sample

Six thousand one hundred and six individuals who responded to the questions on diabetes were included in the analysis. Table 1 shows the characteristics of the study population. The sample comprised of equivalent proportion of male (49.5%) and females (50.5%). Majority of the sample was aged between 18-49 years (60%), of Chinese ethnicity (75.7%), married (59.8%), with university education (29.4%), employed (72%) and with a monthly household income of SGD 4,000 and above (63%).

Prevalence of diabetes and its correlates

The prevalence of diabetes in this population was 9.7% (n= 904). After excluding those with age of onset before 18 years of age, the prevalence of diabetes was 9.5% (n=890). The mean and median age of onset of diabetes was 48.5 (SD= 13.5) and 49 years, respectively. Participants belonging to the age groups of 35-49 years, 50- 64 years and more than 65 years (versus 18-34 years of age), those of Indian ethnicity (versus Chinese), married (versus single), with preuniversity education (versus university), and those who were overweight and obese (versus normal BMI) were more likely to have diabetes. Female gender and those with household income of SGD 6000-9999 (versus less than below SGD 2000) were less likely to be associated with diabetes (Table 1).

#### Diabetes and ACE

Four thousand four hundred and forty-one individuals who responded to the ACE questions were included in the analysis to examine the association between diabetes and types/frequencies of ACEs.

Supplementary Table 1 shows the associations between diabetes and ACEs. In the overall sample, after adjusting for both health risk-behaviors and sociodemographic factors, only those with parental separation, divorce or death of a parent remained significantly associated with diabetes (OR = 1.5, 95% CI = 1.01-2.2).

Examining the interaction effects, revealed a significant interaction between ACE's and age group and diabetes for specific ACES (living with household members who were mentally ill or suicidal, emotional abuse and bullying).

When the sample was stratified by age, we found that the associations of ACEs with diabetes varied considerably across the two age groups (18-49 and >50 years age group). In the younger age group, after controlling for both health risk-behaviors and sociodemographic factors, parental separation, divorce or death of a parent (OR =2.2, 95% CI =1.1-4.4), bullying (OR =5.5, 95% CI =1.4-21.4) and experiencing 3 or more ACEs (OR =2.3, 95% CI =1.1-5.4) remained significantly associated with diabetes. Among older adults, living with household members who were mentally ill or suicidal (OR =0.2, 95% CI =0.1-0.6), and emotional abuse (OR =0.2, 95% CI =0.1-0.5) were less likely to be associated with diabetes after adjusting for health risk-behaviors and sociodemographic factors (Table 2).

Age of onset of diabetes was substantially earlier in the 18-49 years age group (median = 37 years), compared to the age of onset in those belonging to the >50 years age group (median =51 years).

Comorbid depression, resource utilization and health-related quality of life domains among those with diabetes and ACE versus those with diabetes alone

The prevalence of those with both diabetes and ACE was higher than those with diabetes alone (5.7% (n=350) vs 2.1% (n=137)). Table 3 shows the associations between MDD, resource

utilization, and health-related quality of life domains among those with diabetes and ACEs. The prevalence of MDD was significantly higher among those with diabetes and ACEs than those with diabetes alone (3.7% and 0.3% respectively), and after controlling for sociodemographic and health behavior factors in multivariate regression models, those with diabetes and ACEs remained more likely to be associated with increased odds of MDD than those with diabetes only (OR= 19.7, 95% CI = 2.1-187.4).

Having diabetes with ACE was associated with lower scores on several domains of quality of life such as vitality, role emotional and mental health and MCS as compared with diabetes without ACEs. In contrast, diabetes with ACE was associated with higher scores on PCS as compared with diabetes without ACEs.

#### **Discussion**

The current study used nationally representative data from an urban Asian population to examine the association of ACEs with diabetes among those with onset of diabetes after 18 years. Although exposure to any ACE was not associated with increased odds of diabetes, those who had experienced parental separation, death or divorce of a parent, had higher odds of diabetes. In addition, we observed a significant interaction between age and ACEs in relation to odds of diabetes. Among those belonging to the younger age group (< 50 years), parental separation, divorce or death of a parent, bullying and experiencing 3 or more ACEs was significantly associated with diabetes. In contrast, ACEs were not significantly associated with higher odds of diabetes in the older age group.

Associations with specific ACEs vary widely across studies. Monnat and Chandler<sup>46</sup> found that physical abuse, sexual abuse and exposure to domestic violence increased the odds of diabetes in

adult life, while Rich-Edwards et al.<sup>27</sup> found that physical and sexual abuse in childhood and adolescence were associated with risk of type 2 diabetes among adult women. Longitudinal data from the 1979 National Longitudinal Survey of Youth, found that type 2 diabetes was associated with parental death in women<sup>47</sup>. On the other hand, Widom et al.<sup>48</sup> using data from court records found that physical abuse and neglect (largely physical neglect) were associated with diabetes in middle adulthood. Several reasons have been suggested to explain the association of ACEs with the development of diabetes and other chronic physical conditions. ACEs negatively impact neurodevelopment especially that of the hypothalamic–pituitary– adrenal (HPA) axis,<sup>49</sup> cause an elevated level of triglycerides, free fatty acids and glucose in the blood<sup>23</sup> and they result in the elevation of inflammatory markers all of which have been postulated to be involved in the development of chronic illnesses including diabetes.<sup>50</sup>

In the older population i.e., those more than 50 years of age, living with household members who were mentally ill or suicidal and emotional abuse were both less likely to have diabetes. We are currently unable to explain this association. This suggests that while age remains one of the most significant risk factors for diabetes, ACEs are an important risk factor for diabetes among those belonging to the younger age groups. Interestingly the age of onset of diabetes was significantly earlier in the 18 to 49 years age group, compared to the age of onset in those older than 50 years, which raises the possibility that ACEs may be linked to an earlier onset of diabetes and thus is more strongly associated with diabetes in those belonging to the younger age group.

Several studies have highlighted that health risk behaviours which are associated with ACEs may explain poor outcomes such as diabetes in this group. However, upon adjusting for health risk

behaviours, the association between ACEs and diabetes did not change significantly for the overall sample in our study. These results are similar to that of Kreatsoulas et al.<sup>51</sup> who found that when they tested the association of adverse childhood experiences with diabetes through health behaviors which are known risk factors such as obesity, the relationship did not attenuate significantly. The authors suggested that adverse childhood experiences may operate on an independent and potent pathway early in life.

Few studies have examined the impact of ACEs among those with diabetes. Our study in line with our hypothesis found that those with ACE and diabetes were more likely to be associated with MDD, and lower HRQoL. Stress and inflammatory mechanisms have been suggested to explain the association between diabetes and depression,<sup>52</sup> it therefore follows that those with diabetes and ACEs would have greater cumulative stress that could lead to a higher prevalence of depression in this group. On the other hand, a mediation analysis suggested that ACEs were indirectly associated with an increased risk of diabetes in adulthood via depressive symptoms<sup>23</sup>. Corso et al.<sup>32</sup> have reported lower HRQoL among those with ACEs. Thus, ACEs alone or in combination with depression could have led to a lower HRQoL among those with diabetes and ACES.

However, in terms of resource utilization the results did not support our hypothesis. Our study, unlike Hargreaves et al.<sup>53</sup> who reported greater use of emergency rooms and doctor office visits with increasing ACEs as compared to those without any ACEs, did not find increased resource utilisation among those with diabetes and ACEs. Koball et al.<sup>12</sup> examining the impact of ACEs on resource utilization found that patients with higher ACEs were more likely to miss appointments or 'late-cancelled' appointments. Thus, while resource utilization may be low there would be an

enormous cost to the healthcare system. The lack of significant association of those with diabetes and ACEs may thus be due to the fact that we did not examine it in those with high ACEs (current analysis included those with at least one ACE) or it could be that while those with ACEs needed services, they did not attend them.

The prevalence of diabetes in our study sample was 9.7%. Socio-demographic factors and health behaviours including older age, Indian ethnicity as well as being overweight and obese were associated with diabetes, whereas female gender was associated with a lower prevalence of diabetes These findings are similar to the prevalence figures of 9.1% of diabetes in Singapore as reported by the World Health Organisation (WHO) who also identified overweight, obesity and physical inactivity to be significant risk factors.<sup>54</sup> Older age and Indian ethnicity have similarly been identified as risk factors for diabetes in Singapore.<sup>55</sup> These results add support to the validity of the study methodology and our data.

#### Limitations

Some limitations of the study need to be noted so that the gaps can be addressed in future studies. Data on ACEs was captured retrospectively and is there is the possibility of recall bias, and given the sensitivity of the questions there could have been a tendency to under-report the experience. However, we have tried to minimize this by assuring the participants of the confidential nature of the study and that no participant identifiers would be collected. Furthermore, the ACE questionnaire was self-administered by participants to reduce under-reporting as much as possible. Some of the participants refused to complete the ACE questionnaire but the participation rates were comparable to other studies.<sup>41</sup> <sup>56</sup> The study did not assess the severity, age of onset and

duration of ACEs which may have an impact on the development of diabetes.<sup>27 57</sup> Diagnosis of diabetes was also self-reported and thus we would have missed out those who had not been given a clinical diagnosis of diabetes. The questions in our study did not differentiate between Type 1 and Type 2 diabetes. However, Type 1 diabetes onset is rare above the age of 20. The current data included only those with age of onset of diabetes after 18 years and thus, we assume that most cases were those of type 2 diabetes. Lastly, the cross-sectional design limits assertions regarding the sequence of causal events, but this was overcome to a significant extent by excluding those with onset of diabetes before 18 years of age.

#### **Conclusions**

Our study establishes an association between ACEs and diabetes in a multi-ethnic urban population of a developed Asian country. Among those with diabetes, presence of ACEs worsens outcomes in terms of comorbidity with MDD, and poorer HRQoL. Our findings highlight the need to address ACEs across society to improve health outcomes for the population. A multi-agency effort is needed across both health and social care sectors to screen, prevent and provide early care for those who have experienced ACEs. However, we must be cognizant that advancing age and BMI are other important factors associated with diabetes. Thus, efforts to promote regular exercise and healthy lifestyles both in the population and among those with diabetes, must continue along with better identification of, and communication about ACEs within the healthcare setting. We need to develop trauma-informed methods for screening and treating childhood trauma. We also need to create more awareness of both the prevalence and impact of ACEs not just among mental health providers but also among those treating chronic diseases. A collaborative care approach to treat

those with chronic diseases and ACEs that addresses their medical, mental health, and traumarelated issues will improve outcomes for the person and the society as a whole.



#### **Author Statement**

Contributors: SAC, MS and JAV were involved in the planning of the study and provided advice and guidance throughout the conduct study, and manuscript writing. SC, RS, AJ and ES were involved in the conduct of the study. EA conducted the statistical analyses, RVD guided the data analysis. RVD and WLC provided expert opinion. All authors gave critical comment throughout the writing process. All authors were responsible for the conceptualisation and design of the project. All authors approved the final version to be published.

Funding: The study was funded by the Ministry of Health Singapore and Temasek Foundation Innovates.

Competing interests: None declared.

Patient consent for publication: Not required.

**Provenance and peer review:** Not commissioned; externally peer reviewed.

**Ethics approval:** The research was approved by the National Healthcare Group Domain Specific Review Board and given hospital institutional approval.

Data availability statement: Raw data from this study are currently not publicly available but can be made available upon reasonable request to the corresponding author.

#### References

- 1. Felitti VJ, Anda RF, Nordenberg D, *et al.* Relationship of childhood abuse and household dysfunction to many of the leading causes of death in adults. The adverse childhood experiences (ACE) study. *Am J Prev Med* 1998;14(4):245–58.
- 2. Sacks V, Murphey D, Moore K. *Adverse childhood experiences: national and State-level prevalence.* Bethesda, Maryland: Child Trends, 2014.
- 3. Bellis MA, Hughes K, Leckenby N, *et al.* National household survey of adverse childhood experiences and their relationship with resilience to health-harming behaviors in England. *BMC Med* 2014;12:72.
- 4. Nurius P S, Logan-Greene P, Green S. Adverse Childhood Experiences (ACE) within a social disadvantage framework: distinguishing unique, cumulative, and moderated contributions to adult mental health. *J Prev Interv Community* 2012;40:278–90.
- 5. Kelly-Irving M, Lepage B, Dedieu D, *et al*. Childhood adversity as a risk for cancer: findings from the 1958 British birth cohort study. *BMC Public Health* 2013;13:767.
- 6. Su S, Wang X, Pollock JS, *et al.* Adverse childhood experiences and blood pressure trajectories from childhood to young adulthood: the Georgia stress and heart study. *Circulation* 2015;131(19):1674–81.
- 7. Lippard ETC, Nemeroff CB. The Devastating Clinical Consequences of Child Abuse and Neglect: Increased Disease Vulnerability and Poor Treatment Response in Mood Disorders.

  \*\*Am J Psychiatry 2020;177(1):20–36.
- 8. Brown DW, Anda RF, Tiemeier H, *et al.* Adverse childhood experiences and the risk of premature mortality. *Am J Prev Med* 2009;37(5):389–96.

- 9. Lee C, White HR. Effects of childhood maltreatment on violent injuries and premature death during young adulthood among urban high-risk men. *Arch Pediatr Adolesc Med* 2012;166(9):814–20.
- 10. Bellis MA, Hughes K, Hardcastle K, *et al*. The impact of adverse childhood experiences on health service use across the life course using a retrospective cohort study. *J Health Serv Res Policy* 2017;22:168–77.
- 11. Mercado R C, Wiltsey-Stirman, S, Iverson KM. Impact of childhood abuse on physical and mental health Status and health care utilization among female veterans. *Mil Med* 2015;180(10):1065–74.
- 12. Koball AM, Rasmussen C, Olson-Dorff D, *et al*. The relationship between adverse childhood experiences, healthcare utilization, cost of care and medical comorbidities. *Child Abuse Negl* 2019;90:120–6.
- 13. Chartier MJ, Walker JR, Naimark B. Childhood abuse, adult health, and health care utilization: Results from a representative community sample. *Am J Epidemiol* 2007;165(9):1031–38.
- 14. Saeedi P, Petersohn I, Salpea P, *et al*. Global and regional diabetes prevalence estimates for 2019 and projections for 2030 and 2045: Results from the International Diabetes Federation Diabetes Atlas, 9th edition. *Diabetes Res Clin Pract* 2019;107843.
- 15. NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in diabetes since 1980: a pooled analysis of 751 population-based studies with 4.4 million participants. *Lancet* 2016;387(10027):1513–30.
- 16. International Diabetes Federation. *IDF Diabetes Atlas*, 9th ed.; International DiabetesFederation: Brussel, Belgium, 2019; Available online:

- https://diabetesatlas.org/upload/resources/2019/IDF\_Atlas\_9th\_Edition\_2019.pdf (accessed on 20 Feb 2020).
- 17. Huang H, Yan P, Shan Z, *et al.* Adverse childhood experiences and risk of type 2 diabetes: a systematic review and meta-analysis. *Metabolism* 2015;64:1408–18
- 18. Lynch L, Waite R, Davey MP. Adverse childhood experiences and diabetes in adulthood: support for a collaborative approach to primary care. *Contemp Fam Ther* 2013; 35(4):639–55.
- 19. Huffhines L, Noser A, Patton SR. The Link Between Adverse Childhood Experiences and Diabetes. *Curr Diab Rep* 2016;16(6):54.
- 20. Chan O, Inouye K, Riddell MC, *et al.* Diabetes and the hypothalamo-pituitary-adrenal (HPA) axis. *Minerva Endocrinol* 2003;28(2):87–102.
- 21. Rao U, Hammen C, Ortiz LR, *et al.* Effects of early and recent adverse experiences on adrenal response to psychosocial stress in depressed adolescents. *Biol Psychiatry* 2008;64:521–6.
- 22. Diz-Chaves Y, Gil-Lozano M, Toba L, *et al.* Stressing diabetes? The hidden links between insulinotropic peptides and the HPA axis. *J Endocrinol* 2016;230(2):R77–94.
- 23. Deschênes SS, Graham E, Kivimäki M, *et al.* Adverse Childhood Experiences and the Risk of Diabetes: Examining the Roles of Depressive Symptoms and Cardiometabolic Dysregulations in the Whitehall II Cohort Study. *Diabetes Care* 2018;41(10):2120–6.
- 24. Baumeister D, Akhtar R, Ciufolini S, *et al*. Childhood trauma and adulthood inflammation: a meta-analysis of peripheral C-reactive protein, interleukin-6 and tumour necrosis factoralpha. *Mol Psychiatry* 2016;21(5):642–9.

- 25. Yang BZ, Zhang H, Ge W, *et al*. Child abuse and epigenetic mechanisms of disease risk.

  Am *J Prev Med* 2013;44(2):101–7.
- 26. Suderman M, Borghol N, Pappas JJ, *et al*. Childhood abuse is associated with methylation of multiple loci in adult DNA. *BMC Med Genet* 2014;7:1–12.
- 27. Rich-Edwards JW, Spiegelman D, Lividoti Hibert EN, et al. Abuse in childhood and adolescence as a predictor of type 2 diabetes in adult women. Am J Prev Med 2010;39(6):529–36.
- 28. Danese A, Tan M. Childhood maltreatment and obesity: systematic review and metaanalysis. *Mol Psychiatry* 2014;19(5):544–54.
- 29. Davis CR, Dearing E, Usher N, *et al.* Detailed assessments of childhood adversity enhance prediction of central obesity independent of gender, race, adult psychosocial risk and health behaviors. *Metabolism* 2014;63(2):199–206.
- 30. Farr OM, Ko BJ, Joung KE, *et al.* Posttraumatic stress disorder, alone or additively with early life adversity, is associated with obesity and cardiometabolic risk. *Nutr Metab Cardiovasc Dis* 2015;25(5):479–88.
- 31. Danese A, Moffitt TE, Harrington H, *et al.* Adverse childhood experiences and adult risk factors for age-related disease: depression, inflammation, and clustering of metabolic risk markers. *Arch Pediatr Adolesc Med* 2009;163(12):1135–43.
- 32. Corso PS, Edwards VJ, Fang X, *et al.* Health-related quality of life among adults who experienced maltreatment during childhood. *Am J Public Health* 2008;98:1094–100.
- 33. Subramaniam M, Abdin E, Seow E, *et al.* Prevalence, socio-demographic correlates and associations of adverse childhood experiences with mental illnesses: Results from the Singapore Mental Health Study. *Child Abuse Negl* 2020;103:104447.

- 34. Ministry of Health. *National Health Survey*. 2010. Available from from https://www.moh.gov.sg/content/dam/moh\_web/Publications/Reports/2011/NHS2010%20-%20low%20res.pdf Accessed on March 2020.
- 35. Ministry of Health. 2017. *Executive Summary on National Population Health Survey* 2016/17. Available online at https://www.moh.gov.sg/docs/librariesprovider5/resources-statistics/reports/executive-summary-nphs-2016\_17.pdf
- 36. Subramaniam M, Abdin E, Vaingankar JA, *et al.* Tracking the mental health of a nation: prevalence and correlates of mental disorders in the second Singapore mental health study. *Epidemiol Psychiatr Sci Epidemiol Psychiatr Sci* 2020;29:E29.
- 37. Kessler RC, Ustün TB. The World Mental Health (WMH) Survey Initiative Version of the World Health Organization (WHO) Composite International Diagnostic Interview (CIDI). *Int J Methods Psychiatr Res* 2004;13(2):93–121.
- 38. Chong SA, Abdin E, Nan L, *et al.* Prevalence and impact of mental and physical comorbidity in the adult Singapore population. *Ann Acad Med Singapore* 2012;41:105–14.
- 39. World Health Organization. Adverse childhood experiences International questionnaire

  Adverse childhood experiences international questionnaire (ACE-IQ), WHO, Geneva,

  2018. Available online at http://www.who.int/violence\_injury\_prevention/violence
  /activities/adverse childhood experiences/en/
- 40. Bruffaerts R, Demyttenaere K, Borges G, *et al.* Childhood adversities as risk factors for onset and persistence of suicidal behaviour. *Br J Psychiatry* 2010;197(1):20–7.
- 41. Hughes K, Bellis MA, Hardcastle KA, et al. The effect of multiple adverse childhood experiences on health: A systematic review and meta-analysis. Lancet Public Health 2017;2(8):e356–66.

- 42. Scott KM, Von Korff M, Angermeyer MC, *et al.* Association of childhood adversities and early-onset mental disorders with adult-onset chronic physical conditions. *Arch Gen Psychiatry* 2011;68(8):838–44.
- 43. World Health Organization. *WHO Technical Report Series 894; Obesity: Preventing and managing the global epidemic.* 2000. Available from: http://www.who.int/nutrition/publications/obesity/WHO TRS 894/en/
- 44. Ware J Jr, Kosinski M, Keller SD. A 12-item short-form health survey: construction of scales and preliminary tests of reliability and validity. *Med Care* 1996;34(3):220–33.
- 45. Beecham J, Knapp M. *Costing psychiatric interventions*. In: Thornicroft G, Brewin CR, Wing J, editors. Measuring Mental Health Needs. London: Gaskell, 1992:163–83.
- 46. Monnat SM, Chandler RF. Long term physical health consequences of adverse childhood experiences. *Sociol Q* 2015;56(4):723–52.
- 47. Lown EA, Lui CK, Karriker-Jaffe K, *et al.* Adverse childhood events and risk of diabetes onset in the 1979 National longitudinal survey of youth cohort. *BMC Public Health* 2019;19(1):1007.
- 48. Widom CS, Czaja SJ, Bentley T, *et al.* A prospective investigation of physical health outcomes in abused and neglected children: new findings from a 30-year follow-up. *Am J Public Health* 2012;102(6):1135–44.
- 49. Kalmakis KA, Meyer JS, Chiodo L, *et al.* Adverse childhood experiences and chronic hypothalamic-pituitary-adrenal activity. *Stress* 2015;18(4):446–50.

- 50. Heard-Garris N, Davis MM, Estabrook R, *et al.* Adverse childhood experiences and biomarkers of inflammation in a diverse cohort of early school-aged children. *Brain Behav Immun Health* 2020;1:100006.
- 51. Kreatsoulas C, Fleegler EW, Kubzansky LD. Young Adults and Adverse Childhood Events: A Potent Measure of Cardiovascular Risk. *Am J Med* 2019;132(5):605–13.
- 52. Badescu SV, Tataru C, Kobylinska L, *et al*. The association between diabetes mellitus and depression. *J Med Life* 2016;9(2):120–25.
- 53. Hargreaves MK, Mouton CP, Liu J, *et al.* Adverse Childhood Experiences and Health Care Utilization in a Low-Income Population. *J Health Care Poor Underserved* 2019;30(2):749–67.
- 54. World Health Organization. *Global report on diabetes*. Geneva, Switzerland, 2016.
- 55. National Registry of Diseases Office. *Information paper on diabetes in Singapore*. 2011.

  Available online at https://www.nrdo.gov.sg/docs/librariesprovider3/Publications--Kidney-Failure/info paper diabetes 2011nov.pdf
- 56. Hauser W, Schmutzer G, Brahler E, *et al.* Maltreatment in childhood and adolescence: Results from a survey of a representative sample of the German population. *Dtsch Arztebl Int* 2011;108:287–94.
- 57. Karavanaki K, Tsoka E, Liacopoulou M, *et al.* Psychological stress as a factor potentially contributing to the pathogenesis of Type 1 diabetes mellitus. *J Endocrinol Invest* 2008;31(5):406–15.

Table 1. Sociodemographic and lifestyle characteristics of participants and their association with diabetes\*

		N	Weighted %		Model 1			Model 2	
Overall		6126	100.0						
Variables				OR	95%	6 CI	OR	95%	6 CI
Age groups	18-34	1707	30.4	1.0			1.0		
(Age in years)	35-49	1496	29.6	5.9	2.4	14.3	5.1	2.1	12.2
	50-64	1626	26.9	12.5	5.2	30.1	10.9	4.5	26.3
	65+	1297	13.1	20.1	8.2	49.3	19.1	7.6	47.8
Gender	Male	3068	49.6	1.0			1.0		
	Female	3058	50.4	0.6	0.5	0.8	0.6	0.5	0.9
Ethnicity	Chinese	1782	75.7	1.0			1.0		
	Malay	1990	12.5	1.6	1.2	2.1	1.1	0.8	1.5
	Indian	1844	8.7	2.4	1.9	3.2	1.9	1.5	2.6
	Others	510	3.1	0.9	0.5	1.4	0.8	0.5	1.2
Marital status	Single	1544	31.0	1.0			1.0		
	Married	3843	59.8	2.6	1.4	4.8	2.5	1.4	4.7
	Divorced	343	5.2	1.3	0.6	3.0	1.3	0.6	2.8
	Widowed	396	4.1	1.9	0.9	4.0	1.9	0.9	4.0
Education	University	1455	29.4	1.0			1.0		
	Primary	1187	16.3	1.8	1.0	3.1	1.7	0.9	3.1
	Secondary	1648	23.0	1.4	0.8	2.3	1.3	0.8	2.3
	Pre-University	304	6.0	2.8	1.4	5.6	2.7	1.3	5.4
	Vocational	508	6.3	0.8	0.4	1.8	0.8	0.4	1.9
	Polytechnic Diploma	1024	19.0	0.7	0.4	1.3	0.7	0.4	1.3
Employment	Employed	4055	72.0	1.0			1.0		
	Unemployed	354	5.3	1.5	1.1	2.1	1.4	1.0	2.0
	Economically inactive	1716	22.7	1.6	0.8	3.1	1.5	0.7	3.0

Income (SGD)	Below 2000	1139	16.4	1.0			1.0		
	2000-3999	1327	19.9	0.9	0.6	1.3	0.8	0.6	1.3
	4000-5999	1111	21.4	0.8	0.5	1.2	0.7	0.5	1.2
	6000-9999	1001	218	0.5	0.3	0.8	0.4	0.3	0.7
	10000+	860	20.4	0.9	0.5	1.4	0.9	0.5	1.4
Smoking status	Never Smoked	4181	73.4				1.0		
	Current smoker	1176	16.1				0.9	0.5	1.3
	Past Smoker	750	10.5				1.5	1.0	2.2
Body Mass Index (BMI)	Normal	2679	54				1.0		
,	Underweight	304	6.5				0.6	0.2	1.7
	Overweight	1923	28.1				1.5	1.1	2.1
	Obese	1077	11.5				3.8	2.4	5.2
Lifetime Drinking status	Non-drinker	2460	22.7				1.0		
	No binge	2995	63.6				0.9	0.7	1.2
	Binge drinking	671	13.7				0.8	0.5	1.4

<sup>\*</sup>includes those with age of onset after 18 years

Model 1 adjusted for sociodemographic characteristics (age group, gender, ethnicity, marital status, education level, employment and income).

Model 2 adjusted for sociodemographic characteristics (age group, gender, ethnicity, marital status, education level, employment and income) and all health risk behaviors (smoking, lifestyle drinking status and obesity).

OR = odds ratio (multiple logistic regression); CI = confidence interval

Table 2. Association between type and frequencies of adverse childhood experiences (ACE) and diabetes\* after controlling for health risk behaviors and socio-demographic factors

	Overall sample Main effect			18-	18-49 years			50 years and older			
	OR	95%	% CI	OR	95%	6 CI	OR	95%	6 CI		
Emotional neglect	1.3	0.9	1.9	1.4	0.7	2.6	1.2	0.8	1.8		
Physical neglect Living with household members who were	1.0#	0.6	2.0	2.3	0.9	5.9	0.7	0.3	1.5		
substance abusers	1.0	0.5	2.0	0.9	0.3	2.6	1.0	0.4	2.6		
Living with household members who were mentally ill or suicidal Battered mother/female guardian Living with household members who were	0.9# 1.3	0.4 0.6	2.1 2.4	1.6 1.3	0.6 0.6	4.0 3.2	<b>0.2</b> 1.1	0.1 0.4	0.6 2.7		
mprisoned	1.7	0.8	3.6	2.1	0.9	4.7	1.3	0.5	3.4		
Parental separation, divorce or death of a parent	1.5	1.01	2.2	2.2	1.1	4.4	1.3	0.8	1.9		
Emotional abuse	0.8	0.3	2.3	1.3	0.5	3.6	0.2	0.1	0.5		
Physical abuse	1.8#	0.8	4.1	2.4	1.0	6.1	0.9	0.3	2.9		
Sexual abuse	2.3	0.9	5.6	2.4	0.8	7.3	2.2	0.7	7.5		
Bullying	2.9#	0.7	12.8	5.5	1.4	21.4	0.3	0.0	1.4		
Any ACE	1.3	0.9	1.9	1.4	0.7	2.8	1.2	0.7	1.8		
Number of ACEs (Reference=0)											
	1.1	0.7	1.8	1.1	0.5	2.5	1.1	0.7	1.8		
2	1.3	0.8	2.3	1.3	0.5	3.4	1.3	0.7	2.4		
3 and more	1.6	0.9	2.8	2.3	1.1	5.4	1.1	0.6	2.2		

OR= odds ratio; CI = confidence interval.

OR were adjusted for sociodemographic (i.e., age, gender, ethnicity, marital status, education, income) and health risk behaviors (i.e., BMI, drinking and smoking) \*cases with age of onset before 18 years were excluded from the analysis

<sup>#</sup>Significant interaction effect between age group and ACE

Table 3 Associations between major depressive disorder, resource utilization, health-related quality of life domains and those with diabetes and any adverse childhood event

Outcomes		A. Diabetes with ACE (n=350)		B. Diabetes without ACE (n=137)		erence) *		
	n	%	n	%	OR	95% CI		p value
Depression	15	3.7	1	0.3	19.7	2.1	187.4	0.010
Resource utilization								
Primary care doctor	172	44.8	66	37.2	1.1	0.5	2.2	0.855
Restructured hospital doctor	105	27.0	40	26.7	1.3	0.6	2.6	0.513
Private hospital/clinic doctors	107	30.3	36	39.3	0.8	0.3	1.7	0.483
	Mean	SD	Mean	SD	Beta coefficient	95% CI		p value
Number of hospitalisations	0.5	4.2	0.3	3.2	0.2	-0.5	0.8	0.607
Quality of life domains	Mean	SD	Mean	SD	Beta coefficient	95% CI		p value
Physical functioning	51.0	11.8	51.1	11.2	0.4	-2.0	2.9	0.736
Role physical	52.6	9.6	54.8	7.6	-1.8	-3.8	0.2	0.075
Bodily pain	27.3	10.7	24.5	8.5	2.1	-0.3	4.6	0.087
General health	48.2	12.8	45.0	11.9	2.6	-0.7	6.0	0.120
Vitality	53.8	11.8	57.4	10.0	-3.5	-6.2	-0.8	0.014
Social functioning	53.7	9.3	55.4	6.1	-1.3	-3.3	0.7	0.208
Role emotional	53.3	8.9	55.5	4.4	-1.7	-3.0	-0.5	0.011
Mental health	55.1	10.4	58.8	7.2	-4.0	-6.2	-1.8	< 0.001
PCS	42.0	7.2	40.0	5.9	2.0	0.5	3.4	0.009
MCS	57.5	10.6	61.6	6.1	-3.9	-5.8	-1.9	< 0.001

<sup>\*</sup>Regression analyses were conducted after controlling for sociodemographic (i.e., age, gender, ethnicity, marital status, education, income) and health risk behaviors (i.e., BMI, smoking and drinking)

TO COLONIA ONL

Supplementary Table 1a. Association between diabetes\* and type/frequencies of ACEs in the overall sample

		Model 1	Model 2			
	OR	95% CI		OR	95%	% CI
Emotional neglect	1.7	1.3	2.3	1.7	1.2	2.3
Physical neglect	1.7	1.0	2.8	1.5	0.9	2.7
Living with household members who were substance abusers	0.9	0.5	1.8	1.1	0.5	2.1
Living with household members who were mentally ill or suicidal	0.6	0.3	1.4	0.7	0.3	1.5
Battered mother/female guardian	0.9	0.5	1.6	0.9	0.5	1.6
Living with household members who were imprisoned		0.7	2.5	1.2	0.6	2.4
Parental separation, divorce or death of a parent		1.3	2.4	1.7	1.2	2.4
Emotional abuse	0.4	0.2	0.9	0.4	0.2	0.9
Physical abuse	0.9	0.5	2.0	0.9	0.4	2.0
Sexual abuse	1.5	0.7	3.3	1.7	0.7	3.8
Bullying	1.1	0.4	3.5	1.3	0.4	4.1
Any ACE	1.6	1.2	2.3	1.6	1.2	2.3
Number of ACEs (Reference=0)						
1	1.5	1.1	2.3	1.6	1.1	2.3
2	1.9	1.2	2.9	1.9	1.2	2.9
3 and more	1.6	0.9	2.5	1.6	1.0	2.7

Model 1: Crude odds ratio (OR)
Model 2: Adjusted OR after controlling for health risk behaviors (i.e., BMI, drinking and smoking)
\*cases with age of onset before 18 years were excluded from the analysis

Supplementary Table 1b. Associations between diabetes\* and type/frequencies of ACEs stratified by age.

	18-	49 yea	rs		50 year	rs and o	older		18-49	years	50 yea	ırs and	older
			M	odel	odel 1					Mo	odel 2		
	OR	95%	6 CI		OR	95%	6 CI	OR	959	% CI	OR	95%	% CI
Emotional neglect	1.5	0.9	2.8		1.3	0.9	1.9	1.4	0.8	2.7	1.3	0.9	1.8
Physical neglect	3.4	1.3	9.2		0.8	0.4	1.5	2.8	1.0	8.0	0.9	0.5	1.7
Living with household members who													
were substance abusers	1.1	0.3	3.4		1.0	0.4	2.2	1.3	0.4	3.9	1.0	0.4	2.4
Living with household members who													
were mentally ill or suicidal	1.7	0.6	4.8		0.3	0.1	0.6	1.9	0.7	5.4	0.3	0.1	0.6
Battered mother/female guardian	1.5	0.6	3.7		0.9	0.4	2.1	1.4	0.6	3.5	0.8	0.4	1.9
Living with household members who													
were imprisoned	2.0	0.9	4.9		1.1	0.4	3.0	1.8	0.7	4.5	1.0	0.4	2.6
Parental separation, divorce or death of a parent	2.5	1.3	4.6		1.2	0.8	1.8	2.4	1.3	4.4	1.2	0.8	1.8
Emotional abuse	1.0	0.4	2.6		0.2	0.1	0.5	1.0	0.4	2.7	0.2	0.1	0.4
Physical abuse	2.3	0.9	5.8		0.8	0.3	2.2	2.1	0.8	5.4	0.7	0.2	2.2
Sexual abuse	2.6	0.9	7.5		2.1	0.7	6.9	2.8	0.9	8.4	1.8	0.5	6.5
Dullaine			11.										
Bullying	3.2	0.9	0		0.3	0.0	1.3	3.7	1.0	13.0	0.3	0.1	1.4
Any ACE	1.5	0.8	2.8		1.3	0.8	1.9	1.4	0.8	2.8	1.3	0.9	2.0
Number of ACEs (Reference=0)													
1	1.1	0.5	2.4		1.2	0.8	2.0	1.1	0.5	2.5	1.3	0.8	2.1
2	1.3	0.5	3.3		1.4	0.8	2.4	1.2	0.5	3.2	1.4	0.8	2.5
3 and more	2.5	1.1	5.5		1.1	0.6	2.1	2.4	1.1	5.5	1.0	0.5	2.0

Model 1: Crude odds ratio (OR);

Model 2: Adjusted OR after controlling for health risk behaviors (i.e., BMI, drinking and smoking)

STROBE Statement—Checklist of items that should be included in reports of cross-sectional studies

	Item No	Recommendation	Page No
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title	3
		or the abstract	
		(b) Provide in the abstract an informative and balanced summary of	3
		what was done and what was found	
Introduction			•
Background/rationale	2	Explain the scientific background and rationale for the investigation	5-6
		being reported	
Objectives	3	State specific objectives, including any prespecified hypotheses	7
Methods			
Study design	4	Present key elements of study design early in the paper	7-10
Setting	5	Describe the setting, locations, and relevant dates, including periods of	7-8
_		recruitment, exposure, follow-up, and data collection	
Participants	6	(a) Give the eligibility criteria, and the sources and methods of	7-8
-		selection of participants	
Variables	7	Clearly define all outcomes, exposures, predictors, potential	8-10
		confounders, and effect modifiers. Give diagnostic criteria, if	
		applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of	8-10
measurement		methods of assessment (measurement). Describe comparability of	
		assessment methods if there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	NA
Study size	10	Explain how the study size was arrived at	NA
Quantitative variables	11	Explain how quantitative variables were handled in the analyses. If	10
Ç		applicable, describe which groupings were chosen and why	
Statistical methods	12	(a) Describe all statistical methods, including those used to control for	11
		confounding	
		(b) Describe any methods used to examine subgroups and interactions	11
		(c) Explain how missing data were addressed	11
		(d) If applicable, describe analytical methods taking account of	10-11
		sampling strategy	
		(e) Describe any sensitivity analyses	NA
Results		(E) Described any serious sty analyses	1,112
Participants	13*	(a) Report numbers of individuals at each stage of study—eg numbers	12
- marpanto	13	potentially eligible, examined for eligibility, confirmed eligible,	
		included in the study, completing follow-up, and analysed	
		(b) Give reasons for non-participation at each stage	12
		(c) Consider use of a flow diagram	NA
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical,	12-13
Descriptive data	14.	social) and information on exposures and potential confounders	28-31
		(b) Indicate number of participants with missing data for each variable	28-31
		of interest	20-31
		or interest	

Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted	30-31
		estimates and their precision (eg, 95% confidence interval). Make clear	
		which confounders were adjusted for and why they were included	
		(b) Report category boundaries when continuous variables were	30-31
		categorized	
		(c) If relevant, consider translating estimates of relative risk into	NA
		absolute risk for a meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and	13
		interactions, and sensitivity analyses	
Discussion			
Key results	18	Summarise key results with reference to study objectives	14-17
Limitations	19	Discuss limitations of the study, taking into account sources of	17-18
		potential bias or imprecision. Discuss both direction and magnitude of any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering objectives,	18-19
•		limitations, multiplicity of analyses, results from similar studies, and	
		other relevant evidence	
Generalisability	21	Discuss the generalisability (external validity) of the study results	17
Other information			
Funding	22	Give the source of funding and the role of the funders for the present	20
		study and, if applicable, for the original study on which the present	
		article is based	

<sup>\*</sup>Give information separately for exposed and unexposed groups.

**Note:** An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.

## **BMJ Open**

# Association of Adverse Childhood Experiences with Diabetes in Adulthood: Results of a cross-sectional epidemiological survey in Singapore

Journal:	BMJ Open
Manuscript ID	bmjopen-2020-045167.R1
Article Type:	Original research
Date Submitted by the Author:	08-Feb-2021
Complete List of Authors:	Subramaniam, Mythily; Institute of Mental Health, Research Division; National University Singapore Saw Swee Hock School of Public Health, Public Health Translation Abdin, Edimansyah; Institute of Mental Health, Research Division Vaingankar, Janhavi; Institute of Mental Health, Research Division Chang, Sherilyn; Institute of Mental Health, Research Division Sambasivam, Rajeswari; Institute of Mental Health, Research Division Jeyagurunathan, Anitha; Institute of Mental Health, Research Division Seow, Lee Seng Esmond; Institute of Mental Health, Research Division Van Dam, Rob; National University Singapore Saw Swee Hock School of Public Health, Epidemiology Chow, Wai Leng; Government of Singapore Ministry of Health, Epidemiology and Disease Control Chong, Siow Ann; Institute of Mental Health, Research Division
<b>Primary Subject Heading</b> :	Diabetes and endocrinology
Secondary Subject Heading:	Epidemiology
Keywords:	General diabetes < DIABETES & ENDOCRINOLOGY, EPIDEMIOLOGY, Child protection < PAEDIATRICS

SCHOLARONE™ Manuscripts



I, the Submitting Author has the right to grant and does grant on behalf of all authors of the Work (as defined in the below author licence), an exclusive licence and/or a non-exclusive licence for contributions from authors who are: i) UK Crown employees; ii) where BMJ has agreed a CC-BY licence shall apply, and/or iii) in accordance with the terms applicable for US Federal Government officers or employees acting as part of their official duties; on a worldwide, perpetual, irrevocable, royalty-free basis to BMJ Publishing Group Ltd ("BMJ") its licensees and where the relevant Journal is co-owned by BMJ to the co-owners of the Journal, to publish the Work in this journal and any other BMJ products and to exploit all rights, as set out in our licence.

The Submitting Author accepts and understands that any supply made under these terms is made by BMJ to the Submitting Author unless you are acting as an employee on behalf of your employer or a postgraduate student of an affiliated institution which is paying any applicable article publishing charge ("APC") for Open Access articles. Where the Submitting Author wishes to make the Work available on an Open Access basis (and intends to pay the relevant APC), the terms of reuse of such Open Access shall be governed by a Creative Commons licence – details of these licences and which Creative Commons licence will apply to this Work are set out in our licence referred to above.

Other than as permitted in any relevant BMJ Author's Self Archiving Policies, I confirm this Work has not been accepted for publication elsewhere, is not being considered for publication elsewhere and does not duplicate material already published. I confirm all authors consent to publication of this Work and authorise the granting of this licence.

Title: Association of Adverse Childhood Experiences with Diabetes in Adulthood: Results of a cross-sectional epidemiological survey in Singapore

Authors: Mythily Subramaniam,<sup>1,2</sup> Edimansyah Abdin,<sup>1</sup> Janhavi Vaingankar,<sup>1</sup> Sherilyn Chang, <sup>1</sup> Rajeswari Sambasivam, <sup>1</sup> Anitha Jeyagurunathan, <sup>1</sup> Esmond Seow,<sup>1</sup> Rob M. van Dam,<sup>2</sup> Wai Leng Chow,<sup>3</sup> Siow Ann Chong<sup>1</sup>

<sup>1</sup>Research Division

**Institute of Mental Health** 

**Singapore** 

<sup>2</sup> Epidemiology Domain,

Saw Swee Hock School of Public Health

Singapore

<sup>3</sup> Epidemiology and Communicable Diseases Division

**Ministry of Health** 

Singapore

**Correspondence:** 

**Mythily Subramaniam** 

**Director, Research Division** 

**Institute of Mental Health** 

**Buangkok Green Medical Park** 

10 Buangkok View

Singapore 539747

Tel: 65-6389 3633 Fax: 65-6343 7962

Email: Mythily@imh.com.sg

Word count: 4406



#### Abstract

Objectives Several studies have linked diabetes mellitus to adverse childhood experiences (ACE) While a number of studies have examined the association between ACEs and diabetes in Western populations, few have done it in Asian populations. The current study aimed to examine (i) the association between ACEs and diabetes, including the association after age stratification and (ii) the association of comorbid depression, resource utilization and health related quality of life (HRQoL) among those with diabetes and ACEs in Singapore.

**Settings** Participants were surveyed in their homes or any other preferred venue of their choice.

**Participants** 6126 individuals aged 18 years and above randomly selected among Singapore residents.

**Design** Cross-sectional nationwide epidemiological study.

**Results** Exposure to any ACE was not associated with increased odds of diabetes, however those who had experienced parental separation, death or divorce of a parent, had higher odds of diabetes. In addition, we observed significant interaction between age and ACEs in relation to odds of diabetes. ACEs were significantly associated with higher odds of diabetes mainly in the younger age group. The prevalence of major depressive disorder was significantly higher among those with diabetes and ACEs than those with diabetes alone (3.7% and 0.3% respectively).

**Conclusion** Efforts to promote regular exercise and healthy lifestyles both in the population and among those with diabetes, must continue for the prevention and management of diabetes. The findings emphasise the need to create more awareness of both the prevalence and impact of ACEs among those treating chronic diseases.

Key words: Adverse childhood experiences; Diabetes; Depression; Healthcare utilization; Asian

#### **Article Summary**

#### Strengths and limitations of this study

- 1. The study reports on the association of Adverse Childhood Events (ACEs) with diabetes based on data from a large representative sample of the Singapore population.
- 2. Data on ACEs was captured retrospectively and is there is the possibility of recall bias and given the sensitivity of the questions there could have been a tendency to under-report the experience.
- 3. The study did not assess the severity, age of onset and duration of ACEs which may have an impact on the development of diabetes.
- 4. The cross-sectional design limits assertions regarding the sequence of causal events, but this was overcome to a significant extent by excluding those with onset of diabetes before 18 years of age.

#### Introduction

Adverse Childhood Experiences (ACEs) is a broad construct that includes neglect, abuse as well as household dysfunction resulting from parental mental illness, substance abuse, imprisonment, domestic violence and divorce occurring before a child reaches 18 years of age. The prevalence of ACEs varies considerably across studies depending on the population and the methodology. A study from the USA found an overall prevalence of 46% while another study estimated that about 47% of the population of England had experienced at least one such ACE3 ACEs impact health outcomes across the life course. These include emergence of mental and physical illnesses, 5-7 premature death and higher health care utilization. Studies have attributed the higher utilization among those with ACEs to health comorbidities, functional disability, need for complex care coordination and frequent missing of appointments.

Diabetes mellitus (subsequently referred to as diabetes) is a chronic metabolic condition with an estimated global prevalence of 9.3% (463 million people) in 2019, which is projected to increase to 10.2% (578 million) by 2030.<sup>14</sup> The annual direct cost of diabetes—i.e., cost of treating and managing the disease and its complications, was calculated to be International\$ 825 billion in 2014.<sup>15</sup> Risk factors commonly identified for type 1 diabetes include, genetic susceptibility, viral infections, and toxins while type 2 diabetes has been strongly linked with increasing age, increased body mass index (BMI), ethnicity, and family history.<sup>16</sup>

Over the years, studies have linked diabetes to ACEs. A systematic review and metaanalysis suggested that adverse childhood experiences (ACEs) were associated with a significantly elevated risk of type 2 diabetes in adulthood. Participants who had experienced an ACE had a 32% increased risk of developing type 2 diabetes later in life. Additionally, they found a differential effect of the type of ACE; neglect had the strongest effect while physical abuse had the least.<sup>17</sup> Studies have further shown that endorsement of a single ACE may increase risk for diabetes in adulthood by 11%<sup>18</sup> and this risk may increase two-fold when there are more than three ACEs.<sup>19</sup>A meta-analysis by Jakubowski et al.<sup>20</sup> examined the association of cumulative childhood adversity (index of at least 2 ACEs) on cardiometabolic disease (either cardiovascular disease outcomes and metabolic outcomes – diabetes and metabolic syndrome) and it showed a significant effect of cumulative childhood adversity on adult metabolic outcomes.

Suggested pathophysiological mechanisms linking ACEs with diabetes include, effects on the hypothalamic- pituitary axis, <sup>21-23</sup> metabolic dysregulation i.e., elevated level of triglycerides, free fatty acids, glucose, and insulin caused by the chronic stress from ACEs, <sup>24</sup> elevated levels of inflammatory markers, <sup>25</sup> and epigenetic changes. <sup>26</sup> <sup>27</sup> Psycho-social risk factors have also been proposed as a possible mechanism. Those with a history of ACEs are more likely to have health-risk behaviors such as cigarette smoking, alcohol misuse, infrequent physical activity and having higher BMIs<sup>28-30</sup>- all of which can increase the risk of diabetes. Lastly, studies have shown a significant association of ACEs with posttraumatic stress disorder (PTSD) and depression which in turn are strongly associated with the development of obesity, diabetes and cardiovascular diseases. <sup>19</sup> <sup>24</sup> <sup>31</sup> However, much less is known about the correlates of ACEs for adults who have diabetes. Although in general exposure to ACEs is associated with increased burden to physical illness such as depression, <sup>32</sup> and lower health-related quality of life (QOL), <sup>33</sup> it is not known how or if these variables are related to ACEs in patients who have diabetes.

Singapore is a multi-ethnic country located in Southeast Asia. A recent study found that the lifetime prevalence of ACEs in the adult population of Singapore was 63.9%. ACEs were associated with several mental disorders including major depressive disorder (MDD) and alcohol use disorder.<sup>34</sup> However, the study did not examine the association of ACEs with physical

conditions. The prevalence of diabetes in Singapore is higher compared to the global prevalence, with about 1 in 9 Singaporeans (11.3%) suffering from the disease.<sup>35</sup> Diabetes was the seventh leading cause of disability adjusted life years in Singapore in 2017.<sup>36</sup> While a number of studies have examined the association between ACEs and diabetes in Western populations, few have done it in Asian populations. It is also not known whether those with diabetes and ACEs are more likely to be associated with mental disorders or increased resource utilization.

Accordingly, the current study aimed to examine (i) the association between ACEs and diabetes, including the association after age stratification and (ii) the association of comorbid depression, resource utilization and health related quality of life (HRQoL) among those with diabetes and ACEs. The authors hypothesized that experiencing ACEs in childhood would be associated with diabetes, and that the association will be attenuated by health risk behaviors, including obesity, smoking and alcohol use. We also hypothesized that those with diabetes and ACEs would be significantly associated with MDD, increased resource utilization and lower HRQoL.

#### Methods

The current study comprises secondary analysis of data from the Singapore Mental Health Study conducted in 2016 (SMHS 2016). SMHS 2016 was a population-based, psychiatric epidemiological study conducted among Singapore residents. The study has been described in detail in an earlier article.<sup>37</sup> In brief, a probability sample was randomly selected using a disproportionate stratified sampling design from a national registry of Singaporeans and permanent residents. Residents aged 65 and above, Malays and Indians were over-sampled to ensure that an adequate sample size would be achieved to improve the reliability of estimates for the subgroup

analysis. In all, 6,126 respondents were interviewed as part of the study with a response rate of 69.5%.<sup>37</sup>

#### 2.1. Study procedures

Face to face interviews were conducted in English, Chinese or Malay language depending on the preference of the respondents. Respondents comprised those who were 18 years and older, Singapore citizens or Permanent residents, living in Singapore in a community dwelling during the field period. Trained interviewers from a survey research company administered the survey using computer assisted personal interviews (CAPI). The study was approved by the National Healthcare Group's Domain Specific Review Board. Written informed consent was obtained from all participants and parents or legally acceptable representatives of those aged below 21 years.

#### 2.2. Questionnaires

2.2.1. World Health Organisation – Composite International Diagnostic Interview version 3.0 (WHO-CIDI 3.0)

Diagnoses of lifetime MDD, was generated using the CIDI 3.0 algorithm for the Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV) criteria.<sup>38</sup>

A modified version of the CIDI checklist of chronic medical disorders was used to establish the prevalence of eighteen chronic medical conditions including diabetes.<sup>39</sup> The question was read as 'I'm going to read to you a list of health problems some people have. Has a doctor ever told you that you have any of the following...' This was followed by a list of chronic medical conditions which were considered prevalent in Singapore's population. Those who gave a positive answer to the question were routed to the question "How old were you when you were first diagnosed with the condition?"

#### 2.2.2. Adverse Childhood Experiences—International Questionnaire (ACE -IQ)

The World Health Organisation's international questionnaire was used to establish the prevalence of ACEs in the current study.<sup>40</sup> The questions from ACE-IQ included in the current study cover physical, sexual and emotional abuse and neglect by parents or caregivers; as well as family dysfunction. Only one question was asked to assess sexual abuse – "Did someone touch or fondle you in a sexual way when you did not want them to?" All questions about ACEs pertained to the respondents' first 18 years of life, and responses were binary (yes or no) or frequency-based. From these, a dichotomous variable was created to reflect exposure to each ACE type and category (abuse, neglect, household dysfunction, and bullying). Given the sensitive nature of the questions, the questionnaire was self-administered by the respondent using a tablet. For those who were not literate, the items were read out in a neutral manner by the interviewer. Consistent with previous research, <sup>41-43</sup> the number of ACEs that the respondents reported having experienced was summed into an ACE count (range 0 to 11) and categorised into four groups for analysis: 0 ACE, 1 ACE, 2 ACEs and 3 + ACEs.

#### 2.2.3 Health Risk Behaviors

- *Overweight and obesity:* BMI was calculated as weight in kilograms divided by the square of height in meters; weight and height were self-reported by participants. BMI was coded into underweight (< 18.5), normal (18.5–24.9), overweight (25–29.9), and obese (> = 30), consistent with current recommendations.<sup>44</sup>
- *Alcohol use*: Based on questions from the WHO-CIDI 3.0 as well as a series of questions created to assess binge drinking, participants were classified as non-drinkers, drinker but not binge drinker and those with binge drinking.

- *Smoking history:* Participants were asked if they are current smokers, ex-smokers or non-smokers i.e., have never smoked before. They were then classified as never, past, and current smokers.

#### 2.2.3. SF-12 questionnaire

The Short Form (SF)-12 instrument was used to measure the generic HRQoL.<sup>45</sup> It measures eight domains of health: physical function (PF), role-physical (RP), bodily pain (BP), general health (GH), vitality (VT), social function (SF), role-emotion (RE), and mental health (MH). Scores from these domains are used to generate two summary scores - physical component summary (PCS) and mental component summary (MCS). Both summary scores range between 0 and 100, with higher scores indicating better health. Internal consistency reliability for PCS (Cronbach's alpha = 0.81) and MCS (Cronbach's alpha = 0.82) domain scores were high in the current study.

#### 2.2.4 Resource utilization

Information on healthcare resource utilization was obtained from respondents using an adapted version of the Client Service Receipt Inventory (CSRI).<sup>46</sup> The instrument asked whether respondents had accessed specific healthcare resources during the past 3-month period before the time of the interview. This included care from the polyclinic doctor (government primary care doctors), restructured hospital doctors (a public hospital doctor which is wholly owned by the government), private hospital/clinic doctors, and hospitalisations.

#### 2.2.5 Socio-Demographic Questionnaire

Data on sex, age, ethnicity (Chinese, Malay, Indian, and Others), marital status (not married, married, divorced/ separated and widowed), educational level (primary and below, secondary, vocational/ institute of technical education (ITE), pre-university/ junior college, diploma and

university), employment status (employed, unemployed and economically inactive, i.e., students, homemakers and retirees) and monthly household income was collected. Monthly household income was calculated as the average pre-tax income in the past 12 months, of all family members living in the same household.

#### Statistical analysis

Statistical analyses were carried out using the STATA version 13.0. All estimates were analyzed using survey weights to adjust for oversampling, non-response and poststratification according to age and ethnicity distribution between the survey sample and the Singapore adult population in 2014. The results are presented as 'n' which is the frequency of variable observed/ endorsed in the sample and '%' which is the adjusted prevalence in the population. Mean and standard errors were calculated for continuous variables and frequencies and percentages for categorical variables. Multiple logistic regression was used to determine sociodemographic factors associated with diabetes as well as to determine the association of ACEs (specific experiences and counts) with diabetes (model 1), after adjusting for health risk behavior factors (model 2) and adjusting for both health risk behaviors and sociodemographic factors (model 3). In addition, we tested for interaction between age (18-49 years, and 50+ yrs) and different ACEs (yes/no) in relation to diabetes by including multiplicative interaction terms in the multivariable logistic regression models. We chose the cut-off age as 49 years as the median age of onset of diabetes in the sample was 49 years. Since the interaction between age and several ACEs was significant, we described the associations between ACEs and diabetes stratified for age.

Multiple linear and logistic regression models were also used to examine the associations among those with diabetes and any ACE (versus among those with diabetes alone) and MDD,

primary care visits, restructured hospital doctor visits, private hospital doctor visits, and number of hospitalisations after adjusting for health risk behavior factors and sociodemographic factors.

Those with onset of diabetes up to the age of 18 years were excluded from these analyses as ACEs measure adversities experienced until 18 years. All statistically significant differences were evaluated at the 0.05 level using 2-sided tests.

#### Patient and public involvement

Patients or participants of this study were not involved in the design or development of the study.

#### **Results**

Socio-demographic characteristics of the sample

Our analysis on diabetes risk factors included 6106 individuals who responded to the questions on diabetes. Table 1 shows the characteristics of the study population. The sample comprised of equivalent proportion of male (49.5%) and females (50.5%). The majority of the sample was aged between 18-49 years (60.0%), of Chinese ethnicity (75.7%), married (59.8%), with university education (29.4%), employed (72.0%) and with a monthly household income of SGD 4,000 and above (63.0%).

Prevalence of diabetes and its correlates

The prevalence of diabetes in this population was 9.7% (n= 904). After excluding those with age of onset before 18 years of age, the prevalence of diabetes was 9.5% (n=890). The mean and median age of onset of diabetes was 48.5 (SD=13.5) and 49 years, respectively. Older age, Indian ethnicity (versus Chinese), being married (versus single), having pre-university education (versus university), and being overweight and obese (versus normal BMI) were associated with a greater likelihood of having diabetes. Female gender and having a household income of SGD 6000-9999

(versus less than below SGD 2000) were associated with a lower likelihood of having diabetes (Table 1).

#### Diabetes and ACE

Our analysis on ACE and diabetes included 4441 participants who responded to the ACE questions.

Table 2 shows the associations between ACEs and diabetes. In the overall sample, emotional neglect and parental separation divorce or death of a parent, were significantly associated with higher odds of diabetes, while experiencing emotional abuse was associated with lower odds of diabetes. In addition, experiencing one or two ACEs as compared with no ACE was associated with higher odds of diabetes. However, after adjusting for socio-demographic and health risk-behaviors only parental separation, divorce or death of a parent remained significantly associated with higher odds of diabetes (OR =1.5, 95% CI =1.01-2.2). We observed significant interactions between ACE's and age group in relation to diabetes for several ACEs. Specifically, the association between 'living with household members who were mentally ill or suicidal', emotional abuse and bullying and diabetes differed significantly for younger (18-49 years) and older (≥50 years) participants.

Supplementary Table 1 shows the associations between ACEs and diabetes stratified by age without adjusting for confounders, and adjusting for lifestyle factors in Model 1 and Model 2 respectively. Table 3 shows the fully adjusted associations between ACEs and diabetes stratified according to age. In the younger age group parental separation, divorce or death of a parent (OR =2.2, 95% CI =1.1-4.4), and experiencing 3 or more ACEs (OR =2.3, 95% CI =1.1-5.4) were significantly associated with higher odds of diabetes. In addition, the association between physical

neglect and diabetes was borderline significant in the younger age group (OR =2.3, 95% CI =0.9-5.9). In the older age group, living with household members who were mentally ill or suicidal (OR =0.2, 95% CI =0.1-0.6) and emotional abuse (OR =0.2, 95% CI =0.1-0.5) was significantly associated with lower odds of diabetes.

The age of onset of diabetes was substantially earlier in the younger (18-49 years) age group (median = 37 years) than in the older (>50 years) age group (median = 51 years).

Comorbid depression, resource utilization and health-related quality of life domains among those with diabetes and ACE versus those with diabetes alone

The prevalence of those with both diabetes and ACE was higher than those with diabetes alone (5.7% (n=350) vs 2.1% (n=137)). Table 4 shows the associations between MDD, resource utilization, and health-related quality of life domains among those with diabetes and ACEs. The prevalence of MDD was significantly higher among those with diabetes and ACEs than those with diabetes alone (3.7% and 0.3% respectively), and after controlling for sociodemographic and health behavior factors in multivariate regression models, those with diabetes and ACEs remained more likely to be associated with increased odds of MDD than those with diabetes only (OR= 19.7, 95% CI = 2.1-187.4).

Having diabetes with ACE was associated with lower scores on several domains of quality of life such as vitality, role emotional and mental health, and MCS as compared with diabetes without ACEs. In contrast, diabetes with ACE was associated with higher scores on PCS as compared with diabetes without ACEs.

#### **Discussion**

The current study used nationally representative data from an urban Asian population to examine the association of ACEs with diabetes among those with onset of diabetes after 18 years. Although exposure to any ACE was not associated with increased odds of diabetes, those who had experienced parental separation, death or divorce of a parent, had higher odds of diabetes. In addition, we observed a significant interaction between age and ACEs in relation to the odds of diabetes. Among those belonging to the younger age group (< 50 years), parental separation, divorce or death of a parent, bullying and experiencing 3 or more ACEs was significantly associated with diabetes. In contrast, ACEs were not significantly associated with higher odds of diabetes in the older age group.

Associations with specific ACEs vary widely across studies. Monnat and Chandler<sup>47</sup> found that physical abuse, sexual abuse and exposure to domestic violence increased the odds of diabetes in adult life, while Rich-Edwards et al.<sup>28</sup> found that physical and sexual abuse in childhood and adolescence were associated with risk of type 2 diabetes among adult women. Longitudinal data from the 1979 National Longitudinal Survey of Youth, found that type 2 diabetes was associated with parental death in women.<sup>48</sup> On the other hand, Widom et al.<sup>49</sup> using data from court records found that physical abuse and neglect (largely physical neglect) were associated with diabetes in middle adulthood.

Several reasons have been suggested to explain the association of ACEs with the development of diabetes and other chronic physical conditions. ACEs negatively impact neurodevelopment especially that of the hypothalamic–pituitary– adrenal (HPA) axis,<sup>50</sup> cause an elevated level of triglycerides, free fatty acids and glucose in the blood<sup>23</sup> and they result in the elevation of

inflammatory markers all of which have been postulated to be involved in the development of chronic illnesses including diabetes.<sup>51</sup>

In the older population i.e., those more than 50 years of age, living with household members who were mentally ill or suicidal and emotional abuse were both less likely to have diabetes. We are currently unable to explain this association. This suggests that while age remains one of the most significant risk factors for diabetes, ACEs are an important risk factor for diabetes among those belonging to the younger age groups. Interestingly the age of onset of diabetes was significantly earlier in the 18 to 49 years age group, compared to the age of onset in those older than 50 years, which raises the possibility that ACEs may be linked to an earlier onset of diabetes and thus is more strongly associated with diabetes in those belonging to the younger age group. A study examining the association of ACEs with chronic diseases found that young adults who had experienced four or more ACEs (compared to no ACEs) were at a much greater risk of both prediabetes and diabetes. The incident rate ratios (IRR) for diabetes, and prediabetes were highest in the youngest age group and lower in the older age groups. The authors suggested that the attenuated IRR in the older age groups, could be due to the increase in other risk factors which an individual may be exposed to regardless of any history of ACE.<sup>52</sup> Survival bias could also account for the lower association between ACE and diabetes among the older age group in our study, as other studies have observed a reduction in life expectancy among those exposed to a high burden of ACEs.53

Several studies have highlighted that health risk behaviours which are associated with ACEs may explain poor outcomes such as diabetes in this group. However, upon adjusting for health risk

behaviours, the association between ACEs and diabetes did not change significantly for the overall sample in our study. These results are similar to that of Kreatsoulas et al.<sup>54</sup> who found that when they tested the association of adverse childhood experiences with diabetes through health behaviors which are known risk factors such as obesity, the relationship did not attenuate significantly. The authors suggested that adverse childhood experiences may operate on an independent and potent pathway early in life.

Few studies have examined the impact of ACEs among those with diabetes. Our study in line with our hypothesis found that those with ACE and diabetes were more likely to be associated with MDD, and lower HRQoL. Stress and inflammatory mechanisms have been suggested to explain the association between diabetes and depression,<sup>55</sup> it therefore follows that those with diabetes and ACEs would have greater cumulative stress that could lead to a higher prevalence of depression in this group. On the other hand, a mediation analysis suggested that ACEs were indirectly associated with an increased risk of diabetes in adulthood via depressive symptoms.<sup>24</sup> Corso et al.<sup>33</sup> have reported lower HRQoL among those with ACEs. Thus, ACEs alone or in combination with depression could have led to a lower HRQoL among those with diabetes and ACES.

However, in terms of resource utilization the results did not support our hypothesis. Our study, unlike Hargreaves et al.<sup>56</sup> who reported greater use of emergency rooms and doctor office visits with increasing ACEs as compared to those without any ACEs, did not find increased resource utilisation among those with diabetes and ACEs. Koball et al.<sup>12</sup> examining the impact of ACEs on resource utilization found that patients with higher ACEs were more likely to miss appointments or 'late-cancelled' appointments. Thus, while resource utilization may be low there would be an

enormous cost to the healthcare system. The lack of significant association of those with diabetes and ACEs may thus be due to the fact that we did not examine it in those with high ACEs (current analysis included those with at least one ACE) or it could be that while those with ACEs needed services, they did not attend them.

The prevalence of diabetes in our study sample was 9.7%. Socio-demographic factors and health behaviours including older age, Indian ethnicity as well as being overweight and obese were associated with diabetes, whereas female gender was associated with a lower prevalence of diabetes These findings are similar to the prevalence figures of 9.1% of diabetes in Singapore as reported by the World Health Organisation (WHO) who also identified overweight, obesity and physical inactivity to be significant risk factors.<sup>57</sup> Older age and Indian ethnicity have similarly been identified as risk factors for diabetes in Singapore.<sup>58</sup> These results add support to the validity of the study methodology and our data.

#### Limitations

Some limitations of the study need to be noted so that the gaps can be addressed in future studies. Data on ACEs was captured retrospectively and is there is the possibility of recall bias, and given the sensitivity of the questions there could have been a tendency to under-report the experience. However, we have tried to minimize this by assuring the participants of the confidential nature of the study and that no participant identifiers would be collected. Furthermore, the ACE questionnaire was self-administered by participants to reduce under-reporting as much as possible. Some of the participants refused to complete the ACE questionnaire but the participation rates were comparable to other studies.<sup>42</sup> <sup>59</sup> The study did not assess the severity, age of onset and

duration of ACEs which may have an impact on the development of diabetes.<sup>28 60</sup> Diagnosis of diabetes was also self-reported and thus we would have missed out those who had not been given a clinical diagnosis of diabetes. The questions in our study did not differentiate between type 1 and type 2 diabetes. Type 1 diabetes has an autoimmune aetiology and risk factors are different than for type 2 diabetes. Including those with type 1 diabetes could have led to weakening of the association between ACEs and diabetes in our study as ACEs are not likely to be associated with type 1 diabetes. 61 However, onset of Type 1 diabetes is rare in adults. The current analysis only included those with age of onset of diabetes after age 18 years and diabetes is therefore likely to predominantly be type 2 diabetes. Local data has established a high prevalence of diabetes among Indians.<sup>35</sup>However, the survey questionnaire was not available in Tamil which is one of the official languages of Singapore and spoken by a large proportion of Indians. This may have affected the participation of people of Indian ethnicity and the prevalence of diabetes established in the study. However, the authors believe that this would not have affected the study significantly as English literacy is high in this group. A study by Mathew et al. 62 found that 86% of Indians above the age of 65 and 91% of those in the age range of 51-65 years can speak English well or very well. Lastly, the cross-sectional design limits assertions regarding the sequence of causal events, but this was overcome to a significant extent by excluding those with onset of diabetes before 18 years of age.

#### **Conclusions**

Our study establishes an association between ACEs and diabetes in a multi-ethnic urban population of a developed Asian country. Among those with diabetes, presence of ACEs worsens outcomes in terms of comorbidity with MDD, and poorer HRQoL. Our findings highlight the need to address ACEs across society to improve health outcomes for the population. A multi-agency effort is

needed across both health and social care sectors to screen, prevent and provide early care for those who have experienced ACEs. However, we must be cognizant that advancing age and BMI are other important factors associated with diabetes. Thus, efforts to promote regular exercise and healthy lifestyles both in the population and among those with diabetes, must continue along with better identification of, and communication about ACEs within the healthcare setting. We need to develop trauma-informed methods for screening and treating childhood trauma. We also need to create more awareness of both the prevalence and impact of ACEs not just among mental health providers but also among those treating chronic diseases. A collaborative care approach to treat those with chronic diseases and ACEs that addresses their medical, mental health, and trauma-related issues will improve outcomes for the person and the society as a whole.

#### **Author Statement**

Contributors: SAC, MS and JAV were involved in the planning of the study and provided advice and guidance throughout the conduct study, and manuscript writing. SC, RS, AJ and ES were involved in the conduct of the study. EA conducted the statistical analyses. RVD guided the data analysis. RVD and WLC provided expert opinion. All authors gave critical comment throughout the writing process. All authors were responsible for the conceptualisation and design of the project. All authors approved the final version to be published.

**Funding:** The study was funded by the Ministry of Health Singapore and Temasek Foundation Innovates. Grant Number: NA

Competing interests: None declared.

Patient consent for publication: Not required.

**Provenance and peer review:** Not commissioned; externally peer reviewed.

**Ethics approval:** The research was approved by the National Healthcare Group Domain Specific Review Board (2015/01035) and given hospital institutional approval.

**Data availability statement:** Raw data from this study are currently not publicly available but can be made available upon reasonable request to the corresponding author.

#### References

- 1. Felitti VJ, Anda RF, Nordenberg D, *et al.* Relationship of childhood abuse and household dysfunction to many of the leading causes of death in adults. The adverse childhood experiences (ACE) study. *Am J Prev Med* 1998;14(4):245–58.
- 2. Sacks V, Murphey D, Moore K. *Adverse childhood experiences: national and State-level prevalence.* Bethesda, Maryland: Child Trends, 2014.
- 3. Bellis MA, Hughes K, Leckenby N, *et al.* National household survey of adverse childhood experiences and their relationship with resilience to health-harming behaviors in England. *BMC Med* 2014;12:72.
- 4. Nurius P S, Logan-Greene P, Green S. Adverse Childhood Experiences (ACE) within a social disadvantage framework: distinguishing unique, cumulative, and moderated contributions to adult mental health. *J Prev Interv Community* 2012;40:278–90.
- 5. Kelly-Irving M, Lepage B, Dedieu D, *et al*. Childhood adversity as a risk for cancer: findings from the 1958 British birth cohort study. *BMC Public Health* 2013;13:767.
- 6. Su S, Wang X, Pollock JS, *et al.* Adverse childhood experiences and blood pressure trajectories from childhood to young adulthood: the Georgia stress and heart study. *Circulation* 2015;131(19):1674–81.
- 7. Lippard ETC, Nemeroff CB. The Devastating Clinical Consequences of Child Abuse and Neglect: Increased Disease Vulnerability and Poor Treatment Response in Mood Disorders.

  \*\*Am J Psychiatry 2020;177(1):20–36.
- 8. Brown DW, Anda RF, Tiemeier H, *et al*. Adverse childhood experiences and the risk of premature mortality. *Am J Prev Med* 2009;37(5):389–96.

- 9. Lee C, White HR. Effects of childhood maltreatment on violent injuries and premature death during young adulthood among urban high-risk men. *Arch Pediatr Adolesc Med* 2012;166(9):814–20.
- Bellis MA, Hughes K, Hardcastle K, et al. The impact of adverse childhood experiences on health service use across the life course using a retrospective cohort study. J Health Serv Res Policy 2017;22:168–77.
- 11. Mercado R C, Wiltsey-Stirman, S, Iverson KM. Impact of childhood abuse on physical and mental health Status and health care utilization among female veterans. *Mil Med* 2015;180(10):1065–74.
- 12. Koball AM, Rasmussen C, Olson-Dorff D, *et al*. The relationship between adverse childhood experiences, healthcare utilization, cost of care and medical comorbidities. *Child Abuse Negl* 2019;90:120–6.
- 13. Chartier MJ, Walker JR, Naimark B. Childhood abuse, adult health, and health care utilization: Results from a representative community sample. *Am J Epidemiol* 2007;165(9):1031–38.
- 14. Saeedi P, Petersohn I, Salpea P, et al. Global and regional diabetes prevalence estimates for 2019 and projections for 2030 and 2045: Results from the International Diabetes Federation Diabetes Atlas, 9th edition. *Diabetes Res Clin Pract* 2019;107843.
- NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in diabetes since 1980: a pooled analysis of 751 population-based studies with 4.4 million participants. *Lancet* 2016;387(10027):1513–30.
- International Diabetes Federation. *IDF Diabetes Atlas*, 9th ed.; International DiabetesFederation: Brussel, Belgium, 2019; Available online:

- https://diabetesatlas.org/upload/resources/2019/IDF\_Atlas\_9th\_Edition\_2019.pdf (accessed on 20 Feb 2020).
- 17. Huang H, Yan P, Shan Z, *et al*. Adverse childhood experiences and risk of type 2 diabetes: a systematic review and meta-analysis. *Metabolism* 2015;64:1408–18
- 18. Lynch L, Waite R, Davey MP. Adverse childhood experiences and diabetes in adulthood: support for a collaborative approach to primary care. *Contemp Fam Ther* 2013; 35(4):639–55.
- 19. Huffhines L, Noser A, Patton SR. The Link Between Adverse Childhood Experiences and Diabetes. *Curr Diab Rep* 2016;16(6):54.
- 20. Jakubowski KP, Cundiff JM, Matthews KA. Cumulative childhood adversity and adult cardiometabolic disease: A meta-analysis. *Health Psychol* 2018;37(8):701–15.
- 21. Chan O, Inouye K, Riddell MC, *et al*. Diabetes and the hypothalamo-pituitary-adrenal (HPA) axis. *Minerva Endocrinol* 2003;28(2):87–102.
- 22. Rao U, Hammen C, Ortiz LR, *et al.* Effects of early and recent adverse experiences on adrenal response to psychosocial stress in depressed adolescents. *Biol Psychiatry* 2008;64:521–6.
- 23. Diz-Chaves Y, Gil-Lozano M, Toba L, *et al.* Stressing diabetes? The hidden links between insulinotropic peptides and the HPA axis. *J Endocrinol* 2016;230(2):R77–94.
- 24. Deschênes SS, Graham E, Kivimäki M, *et al.* Adverse Childhood Experiences and the Risk of Diabetes: Examining the Roles of Depressive Symptoms and Cardiometabolic Dysregulations in the Whitehall II Cohort Study. *Diabetes Care* 2018;41(10):2120–6.

- 25. Baumeister D, Akhtar R, Ciufolini S, *et al*. Childhood trauma and adulthood inflammation: a meta-analysis of peripheral C-reactive protein, interleukin-6 and tumour necrosis factoralpha. *Mol Psychiatry* 2016;21(5):642–9.
- 26. Yang BZ, Zhang H, Ge W, *et al*. Child abuse and epigenetic mechanisms of disease risk.

  Am *J Prev Med* 2013;44(2):101–7.
- 27. Suderman M, Borghol N, Pappas JJ, *et al*. Childhood abuse is associated with methylation of multiple loci in adult DNA. *BMC Med Genet* 2014;7:1–12.
- 28. Rich-Edwards JW, Spiegelman D, Lividoti Hibert EN, *et al.* Abuse in childhood and adolescence as a predictor of type 2 diabetes in adult women. *Am J Prev Med* 2010;39(6):529–36.
- 29. Danese A, Tan M. Childhood maltreatment and obesity: systematic review and metaanalysis. *Mol Psychiatry* 2014;19(5):544–54.
- 30. Davis CR, Dearing E, Usher N, *et al.* Detailed assessments of childhood adversity enhance prediction of central obesity independent of gender, race, adult psychosocial risk and health behaviors. *Metabolism* 2014;63(2):199–206.
- 31. Farr OM, Ko BJ, Joung KE, *et al.* Posttraumatic stress disorder, alone or additively with early life adversity, is associated with obesity and cardiometabolic risk. *Nutr Metab Cardiovasc Dis* 2015;25(5):479–88.
- 32. Danese A, Moffitt TE, Harrington H, *et al.* Adverse childhood experiences and adult risk factors for age-related disease: depression, inflammation, and clustering of metabolic risk markers. *Arch Pediatr Adolesc Med* 2009;163(12):1135–43.
- 33. Corso PS, Edwards VJ, Fang X, *et al.* Health-related quality of life among adults who experienced maltreatment during childhood. *Am J Public Health* 2008;98:1094–100.

- 34. Subramaniam M, Abdin E, Seow E, *et al.* Prevalence, socio-demographic correlates and associations of adverse childhood experiences with mental illnesses: Results from the Singapore Mental Health Study. *Child Abuse Negl* 2020;103:104447.
- 35. Ministry of Health. *National Health Survey*. 2010. Available from from https://www.moh.gov.sg/content/dam/moh\_web/Publications/Reports/2011/NHS2010%20-%20low%20res.pdf Accessed on March 2020.
- 36. Ministry of Health. 2017. Executive Summary on National Population Health Survey 2016/17. Available online at https://www.moh.gov.sg/docs/librariesprovider5/resources-statistics/reports/executive-summary-nphs-2016 17.pdf
- 37. Subramaniam M, Abdin E, Vaingankar JA, *et al.* Tracking the mental health of a nation: prevalence and correlates of mental disorders in the second Singapore mental health study. *Epidemiol Psychiatr Sci Epidemiol Psychiatr Sci* 2020;29:E29.
- 38. Kessler RC, Ustün TB. The World Mental Health (WMH) Survey Initiative Version of the World Health Organization (WHO) Composite International Diagnostic Interview (CIDI). *Int J Methods Psychiatr Res* 2004;13(2):93–121.
- 39. Chong SA, Abdin E, Nan L, *et al.* Prevalence and impact of mental and physical comorbidity in the adult Singapore population. *Ann Acad Med Singapore* 2012;41:105–14.
- 40. World Health Organization. *Adverse childhood experiences International questionnaire*\*Adverse childhood experiences international questionnaire (ACE-IQ), WHO, Geneva,

  2018. Available online at http://www.who.int/violence\_injury\_prevention/violence

  /activities/adverse\_childhood\_experiences/en/
- 41. Bruffaerts R, Demyttenaere K, Borges G, *et al.* Childhood adversities as risk factors for onset and persistence of suicidal behaviour. *Br J Psychiatry* 2010;197(1):20–7.

- 42. Hughes K, Bellis MA, Hardcastle KA, *et al.* The effect of multiple adverse childhood experiences on health: A systematic review and meta-analysis. *Lancet Public Health* 2017;2(8):e356–66.
- 43. Scott KM, Von Korff M, Angermeyer MC, *et al.* Association of childhood adversities and early-onset mental disorders with adult-onset chronic physical conditions. *Arch Gen Psychiatry* 2011;68(8):838–44.
- 44. World Health Organization. *WHO Technical Report Series 894; Obesity: Preventing and managing the global epidemic.* 2000. Available from: http://www.who.int/nutrition/publications/obesity/WHO TRS 894/en/
- 45. Ware J Jr, Kosinski M, Keller SD. A 12-item short-form health survey: construction of scales and preliminary tests of reliability and validity. *Med Care* 1996;34(3):220–33.
- 46. Beecham J, Knapp M. *Costing psychiatric interventions*. In: Thornicroft G, Brewin CR, Wing J, editors. Measuring Mental Health Needs. London: Gaskell, 1992:163–83.
- 47. Monnat SM, Chandler RF. Long term physical health consequences of adverse childhood experiences. *Sociol Q* 2015;56(4):723–52.
- 48. Lown EA, Lui CK, Karriker-Jaffe K, *et al.* Adverse childhood events and risk of diabetes onset in the 1979 National longitudinal survey of youth cohort. *BMC Public Health* 2019;19(1):1007.
- 49. Widom CS, Czaja SJ, Bentley T, *et al.* A prospective investigation of physical health outcomes in abused and neglected children: new findings from a 30-year follow-up. *Am J Public Health* 2012;102(6):1135–44.
- 50. Kalmakis KA, Meyer JS, Chiodo L, *et al.* Adverse childhood experiences and chronic hypothalamic-pituitary-adrenal activity. *Stress* 2015;18(4):446–50.

- 51. Heard-Garris N, Davis MM, Estabrook R, *et al.* Adverse childhood experiences and biomarkers of inflammation in a diverse cohort of early school-aged children. *Brain Behav Immun Health* 2020;1:100006.
- 52. Sonu S, Post S, Feinglass J. Adverse childhood experiences and the onset of chronic disease in young adulthood. *Prev Med* 2019;123:163–70.
- 53. Brown DW, Anda RF, Tiemeier H, *et al*. Adverse childhood experiences and the risk of premature mortality. *Am J Prev Med* 2009;37(5):389–96.
- 54. Kreatsoulas C, Fleegler EW, Kubzansky LD. Young Adults and Adverse Childhood Events: A Potent Measure of Cardiovascular Risk. *Am J Med* 2019;132(5):605–13.
- 55. Badescu SV, Tataru C, Kobylinska L, *et al*. The association between diabetes mellitus and depression. *J Med Life* 2016;9(2):120–25.
- Hargreaves MK, Mouton CP, Liu J, et al. Adverse Childhood Experiences and Health Care
   Utilization in a Low-Income Population. J Health Care Poor Underserved 2019;30(2):749–67.
- 57. World Health Organization. *Global report on diabetes*. Geneva, Switzerland, 2016.
- 58. National Registry of Diseases Office. *Information paper on diabetes in Singapore*. 2011.

  Available online at https://www.nrdo.gov.sg/docs/librariesprovider3/Publications--Kidney-Failure/info\_paper\_diabetes\_2011nov.pdf
- 59. Hauser W, Schmutzer G, Brahler E, *et al.* Maltreatment in childhood and adolescence: Results from a survey of a representative sample of the German population. *Dtsch Arztebl Int* 2011;108:287–94.

- 60. Karavanaki K, Tsoka E, Liacopoulou M, *et al.* Psychological stress as a factor potentially contributing to the pathogenesis of Type 1 diabetes mellitus. *J Endocrinol Invest* 2008;31(5):406–15.
- 61. Bengtsson J, Byberg S, Carstensen B, *et al.* Accumulation of childhood adversities and type 1 diabetes risk: a register-based cohort study of all children born in Denmark between 1980 and 2015. *Int J Epidemiol* 2020;49(5):1604–13.
- 62. Mathew M, Tay M, Selvarajen S, Tan ZH. *Language proficiency, identity & management:* results from the IPS survey on race, religion & language. IPS Exchange, Number 15 June 2020. Available online at https://lkyspp.nus.edu.sg/docs/default-source/ips/ips-exchange-series-15.pdf Last accessed on 30 Jan 2021

Table 1. Sociodemographic and lifestyle characteristics of participants and their association with diabetes\*

		N	Weighted %	M	odel 1	Me	odel 2
Overall		6106	100.0				
Variables				OR	95% CI	OR	95% CI
Age groups	18-34	1706	30.5	1.0		1.0	
(Age in years)	35-49	1493	29.6	5.9	2.4-14.3	5.1	2.1-12.2
	50-64	1617	26.8	12.5	5.2-30.1	10.9	4.5-26.3
	65+	1290	13.1	20.1	8.2-49.3	19.1	7.6-47.8
Gender	Male	3057	49.5	1.0		1.0	
	Female	3049	50.5	0.6	0.5-0.8	0.6	0.5-0.9
Ethnicity	Chinese	1776	75.7	1.0		1.0	
	Malay	1980	12.5	1.6	1.2-2.1	1.1	0.8-1.5
	Indian	1841	8.7	2.4	1.9-3.2	1.9	1.5-2.6
	Others	509	3.1	0.9	0.5-1.4	0.8	0.5-1.2
Marital status	Single	1542	31.0	1.0		1.0	
	Married	3830	59.8	2.6	1.4-4.8	2.5	1.4-4.7
	Divorced	342	5.2	1.3	0.6-3.0	1.3	0.6-2.8
	Widowed	392	4.0	1.9	0.9-4.0	1.9	0.9-4.0
Education	University	1451	29.4	1.0		1.0	
	Primary	1184	16.3	1.8	1.0-3.1	1.7	0.9-3.1
	Secondary	1636	22.9	1.4	0.8-2.3	1.3	0.8-2.3
	Pre-University	304	6.1	2.8	1.4-5.6	2.7	1.3-5.4
	Vocational	507	6.3	0.8	0.4-1.8	0.8	0.4-1.9
	Polytechnic Diploma	1024	19.0	0.7	0.4-1.3	0.7	0.4-1.3
Employment	Employed	4047	72.0	1.0		1.0	
	Unemployed	350	5.2	1.5	1.1-2.1	1.4	1.0-2.0
	Economically inactive	1708	22.8	1.6	0.8-3.1	1.5	0.7-3.0
Income (SGD)	Below 2000	1139	16.4	1.0		1.0	
	2000-3999	1327	19.9	0.9	0.6-1.3	0.8	0.6-1.3
	4000-5999	1111	21.4	0.8	0.5-1.2	0.7	0.5-1.2

	6000-9999	1001	21.8	0.5	0.3-0.8	0.4	0.3-0.7
	10000+	860	20.4	0.9	0.5-1.4	0.9	0.5-1.4
Smoking status	Never Smoked	4175	73.4			1.0	
	Current smoker	1171	16.0			0.9	0.5-1.3
	Past Smoker	750	10.6			1.5	1.0-2.2
	N. I	2679	<i>51</i> 1			1.0	
Body Mass Index (BMI)	Normal	2678	54.1			1.0	
	Underweight	303	6.4			0.6	0.2-1.7
	Overweight	1918	28.1			1.5	1.1-2.1
	Obese	1073	11.4			3.8	2.4-5.2
Lifetime Drinking status	Non-drinker	2447	22.6			1.0	
	No binge	2989	63.7			0.9	0.7-1.2
	Binge drinking	670	13.7			0.8	0.5-1.4

<sup>\*</sup> Cases with age of onset before 18 years were excluded from the analysis

Model 1 adjusted for sociodemographic characteristics (age group, gender, ethnicity, marital status, education level, employment and income).

Model 2 adjusted for sociodemographic characteristics (age group, gender, ethnicity, marital status, education level, employment and income) and all health risk behaviors (smoking, lifestyle drinking status and obesity).

OR = odds ratio (multiple logistic regression); CI = confidence interval Significant associations are highlighted in bold

Page 33 of 37

**BMJ** Open

Table 2. Association between type/frequencies of ACEs and diabetes\* in the overall sample

	N	Model 1	M	odel 2	N	Iodel 3
	OR	95% CI	OR	95% CI	OR	95% CI
Emotional neglect	1.7	1.3-2.3	1.7	1.2-2.3	1.3	0.9-1.9
Physical neglect	1.7	1.0-2.8	1.5	0.9-2.7	$1.0^{\#}$	0.6-2.0
Living with household members who were substance abusers	0.9	0.5-1.8	1.1	0.5-2.1	1.0	0.5-2.0
Living with household members who were mentally ill or suicidal	0.6	0.3-1.4	0.7	0.3-1.5	$0.9^{\#}$	0.4-2.1
Battered mother/female guardian	0.9	0.5-1.6	0.9	0.5-1.6	1.3	0.6-2.4
Living with household members who were imprisoned	1.3	0.7-2.5	1.2	0.6-2.4	1.7	0.8-3.6
Parental separation, divorce or death of a parent	1.8	1.3-2.4	1.7	1.2-2.4	1.5	1.01-2.2
Emotional abuse	0.4	0.2-0.9	0.4	0.2-0.9	0.8	0.3-2.3
Physical abuse	0.9	0.5-2.0	0.9	0.4-2.0	$1.8^{\#}$	0.8-4.1
Sexual abuse	1.5	0.7-3.3	1.7	0.7-3.8	2.3	0.9-5.6
Bullying	1.1	0.4-3.5	1.3	0.4-4.1	$2.9^{\#}$	0.7-12.8
Any ACE	1.6	1.2-2.3	1.6	1.2-2.3	1.3	0.9-1.9
Number of ACEs (Reference=0)						
	1.5	1.1-2.3	1.6	1.1-2.3	1.1	0.7-1.8
2	1.9	1.2-2.9	1.9	1.2-2.9	1.3	0.8-2.3
3 and more	1.6	0.9-2.5	1.6	1.0-2.7	1.6	0.9-2.8

<sup>\*</sup> Cases with age of onset before 18 years were excluded from the analysis

Model 1: Crude odds ratio (OR)

Model 2: OR were adjusted for health risk behaviors (i.e., BMI, drinking and smoking)

Model 3: OR were adjusted for sociodemographic (i.e., age, gender, ethnicity, marital status, education, income) and health risk behaviors (i.e., BMI, drinking and smoking)

Significant associations are highlighted in bold

<sup>\*</sup>Significant interaction effect between age group and ACE

Table 3. Association between type and frequencies of adverse childhood experiences (ACE) and diabetes\* after controlling for health risk behaviors and socio-demographic factors

	18-	-49 years	50 yea	ars and older
	OR	95% CI	OR	95% CI
Emotional neglect	1.4	0.7-2.6	1.2	0.8-1.8
Physical neglect	2.3	0.9-5.9	0.7	0.3-1.5
Living with household members who were substance abusers	0.9	0.3-2.6	1.0	0.4-2.6
Living with household members who were mentally ill or suicidal	1.6	0.6-4.0	0.2	0.1-0.6
Battered mother/female guardian	1.3	0.6-3.2	1.1	0.4-2.7
Living with household members who were imprisoned	2.1	0.9-4.7	1.3	0.5-3.4
Parental separation, divorce or death of a parent	2.2	1.1-4.4	1.3	0.8-1.9
Emotional abuse	1.3	0.5-3.6	0.2	0.1-0.5
Physical abuse	2.4	1.0-6.1	0.9	0.3-2.9
Sexual abuse	2.4	0.8-7.3	2.2	0.7-7.5
Bullying	5.5	1.4-21.4	0.3	0.0-1.4
Any ACE	1.4	0.7-2.8	1.2	0.7-1.8
Number of ACEs (Reference=0)				
1	1.1	0.5-2.5	1.1	0.7-1.8
2	1.3	0.5-3.4	1.3	0.7-2.4
3 and more	2.3	1.1-5.4	1.1	0.6-2.2

<sup>\*</sup>Cases with age of onset before 18 years were excluded from the analysis

OR= odds ratio; CI = confidence interval.

OR were adjusted for sociodemographic (i.e., age, gender, ethnicity, marital status, education, income) and health risk behaviors (i.e., BMI, drinking and smoking) Significant associations are highlighted in bold

Table 4 Associations between combinations of diabetes and adverse childhood events with major depressive disorder, resource utilization, health-related quality of life domains

	A. Diabetes with ACE	B. Diabetes without ACE			
	(n=350)	(n=137)	A vs. B (refe	erence) *	
	n (%)	n (%)	OR	95% CI	p value
Depression	15 (3.7)	1 (0.3)	19.7	2.1-187.4	0.010
Resource utilization					
Primary care doctor	172 (44.8)	66 (37.2)	1.1	0.5-2.2	0.855
Restructured hospital doctor	105 (27.0)	40 (26.7)	1.3	0.6-2.6	0.513
Private hospital/clinic					
doctors	107 (30.3)	36 (39.3)	0.8	0.3-1.7	0.483
				0=0/ 67	
	Mean (SD)	Mean (SD)	Beta coefficient	95% CI	p value
Number of hospitalisations	0.5 (4.2)	0.3 (3.2)	0.2	-0.5-0.8	0.607
Quality of life domains	Mean (SD)	Mean (SD)	Beta coefficient	95% CI	p value
Physical functioning	51.0 (11.8)	51.1 (11.2)	0.4	-2.0-2.9	0.736
Role physical	52.6 (9.6)	54.8 (7.6)	-1.8	-3.8-0.2	0.075
Bodily pain	27.3 (10.7)	24.5 (8.5)	2.1	-0.3-4.6	0.087
General health	48.2 (12.8)	45.0 (11.9)	2.6	-0.7-6.0	0.120
Vitality	53.8 (11.8)	57.4 (10.0)	-3.5	-6.20.8	0.014
Social functioning	53.7 (9.3)	55.4 (6.1)	-1.3	-3.3-0.7	0.208
Role emotional	53.3 (8.9)	55.5 (4.4)	-1.7	-3.00.5	0.011
Mental health	55.1 (10.4)	58.8 (7.2)	-4.0	-6.21.8	< 0.001
PCS	42.0 (7.2)	40.0 (5.9)	2.0	0.5-3.4	0.009
MCS	57.5 (10.6)	61.6 (6.1)	-3.9	-5.81.9	< 0.001

<sup>\*</sup>Regression analyses were conducted after controlling for sociodemographic (i.e., age, gender, ethnicity, marital status, education, income) and health risk behaviors (i.e., BMI, smoking and drinking)

Significant associations are highlighted in bold

### Supplementary Table 1. Associations between diabetes\* and type/frequencies of ACEs stratified by age.

	18-	49 years	50 years	and older	18-	49 years	50 years	and older
		Model 1				Mo		
	OR	95% CI	OR	95% CI	OR	95% CI	OR	95% CI
Emotional neglect	1.5	0.9-2.8	1.3	0.9-1.9	1.4	0.8-2.7	1.3	0.9-1.8
Physical neglect	3.4	1.3-9.2	0.8	0.4-1.5	2.8	1.0-8.0	0.9	0.5-1.7
Living with household members who were substance abusers Living with household members who	1.1	0.3-3.4	1.0	0.4-2.2	1.3	0.4-3.9	1.0	0.4-2.4
were mentally ill or suicidal	1.7	0.6-4.8	0.3	0.1-0.6	1.9	0.7-5.4	0.3	0.1-0.6
Battered mother/female guardian	1.5	0.6-3.7	0.9	0.4-2.1	1.4	0.6-3.5	0.8	0.4-1.9
Living with household members who were imprisoned	2.0	0.9-4.9	1.1	0.4-3.0	1.8	0.7-4.5	1.0	0.4-2.6
Parental separation, divorce or death of a parent	2.5	1.3-4.6	1.2	0.8-1.8	2.4	1.3-4.4	1.2	0.8-1.8
Emotional abuse	1.0	0.4-2.6	0.2	0.1-0.5	1.0	0.4-2.7	0.2	0.1-0.4
Physical abuse	2.3	0.9-5.8	0.8	0.3-2.2	2.1	0.8-5.4	0.7	0.2-2.2
Sexual abuse	2.6	0.9-7.5	2.1	0.7-6.9	2.8	0.9-8.4	1.8	0.5-6.5
Bullying	3.2	0.9-11.0	0.3	0.0-1.3	3.7	1.0-13.0	0.3	0.1-1.4
Any ACE	1.5	0.8-2.8	1.3	0.8-1.9	1.4	0.8-2.8	1.3	0.9-2.0
Number of ACEs (Reference=0)								
1	1.1	0.5-2.4	1.2	0.8-2.0	1.1	0.5-2.5	1.3	0.8-2.1
2	1.3	0.5-3.3	1.4	0.8-2.4	1.2	0.5-3.2	1.4	0.8-2.5
3 and more	2.5	1.1-5.5	1.1	0.6-2.1	2.4	1.1-5.5	1.0	0.5-2.0

<sup>\*</sup>Cases with age of onset before 18 years were excluded from the analysis

Model 1: Crude odds ratio (OR);

Model 2: Adjusted OR after controlling for health risk behaviors (i.e., BMI, drinking and smoking)

Significant associations are highlighted in bold

STROBE Statement—Checklist of items that should be included in reports of cross-sectional studies

	Item No	Recommendation	Page No
Title and abstract	1	(a) Indicate the study's design with a commonly used term in the title or the abstract	3
		(b) Provide in the abstract an informative and balanced summary of	3
		what was done and what was found	
Introduction			
Background/rationale	2	Explain the scientific background and rationale for the investigation being reported	5-6
Objectives	3	State specific objectives, including any prespecified hypotheses	7
Methods		and approximation of the second of the secon	
Study design	4	Present key elements of study design early in the paper	7-10
Setting Setting	5	Describe the setting, locations, and relevant dates, including periods of	7-8
Setting		recruitment, exposure, follow-up, and data collection	/-0
Participants	6	(a) Give the eligibility criteria, and the sources and methods of	7-8
r articipants	O	selection of participants	/-0
Variables	7	Clearly define all outcomes, exposures, predictors, potential	8-10
variables	,	confounders, and effect modifiers. Give diagnostic criteria, if	0-10
		applicable	
Data sources/	8*	For each variable of interest, give sources of data and details of	8-10
	8	methods of assessment (measurement). Describe comparability of	0-10
measurement		assessment methods if there is more than one group	
Bias	9	Describe any efforts to address potential sources of bias	NA
Study size	10	Explain how the study size was arrived at	NA NA
Quantitative variables	11	Explain how the study size was arrived at  Explain how quantitative variables were handled in the analyses. If	10
Qualititative variables	11	applicable, describe which groupings were chosen and why	10
Statistical methods	12	(a) Describe all statistical methods, including those used to control for	11
Statistical methods	12	confounding	11
		(b) Describe any methods used to examine subgroups and interactions	11
		(c) Explain how missing data were addressed	11
		(d) If applicable, describe analytical methods taking account of	10-11
		sampling strategy	10-11
		(e) Describe any sensitivity analyses	NA
D 14		(c) Describe any sensitivity analyses	1 17 1
Results	13*	(a) Report numbers of individuals at each stage of study—eg numbers	12
Participants	13.	potentially eligible, examined for eligibility, confirmed eligible,	12
		included in the study, completing follow-up, and analysed	
			12
		(b) Give reasons for non-participation at each stage	1
Dogarintiza data	1 //*	(c) Consider use of a flow diagram	NA
Descriptive data	14*	(a) Give characteristics of study participants (eg demographic, clinical,	12-13
		social) and information on exposures and potential confounders  (b) Indicate number of participants with missing data for each variable.	28-31
		(b) Indicate number of participants with missing data for each variable of interest	28-31
Outaama data	15*		20 21
Outcome data	15*	Report numbers of outcome events or summary measures	28-31

Main results	16	(a) Give unadjusted estimates and, if applicable, confounder-adjusted	30-31
		estimates and their precision (eg, 95% confidence interval). Make clear	
		which confounders were adjusted for and why they were included	
		(b) Report category boundaries when continuous variables were	30-31
		categorized	
		(c) If relevant, consider translating estimates of relative risk into	NA
		absolute risk for a meaningful time period	
Other analyses	17	Report other analyses done—eg analyses of subgroups and	13
		interactions, and sensitivity analyses	
Discussion			
Key results	18	Summarise key results with reference to study objectives	14-17
Limitations	19	Discuss limitations of the study, taking into account sources of	17-18
		potential bias or imprecision. Discuss both direction and magnitude of	
		any potential bias	
Interpretation	20	Give a cautious overall interpretation of results considering objectives,	18-19
		limitations, multiplicity of analyses, results from similar studies, and	
		other relevant evidence	
Generalisability	21	Discuss the generalisability (external validity) of the study results	17
Other information			
Funding	22	Give the source of funding and the role of the funders for the present	20
		study and, if applicable, for the original study on which the present	
		article is based	

<sup>\*</sup>Give information separately for exposed and unexposed groups.

**Note:** An Explanation and Elaboration article discusses each checklist item and gives methodological background and published examples of transparent reporting. The STROBE checklist is best used in conjunction with this article (freely available on the Web sites of PLoS Medicine at http://www.plosmedicine.org/, Annals of Internal Medicine at http://www.annals.org/, and Epidemiology at http://www.epidem.com/). Information on the STROBE Initiative is available at www.strobe-statement.org.